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# PROVISIONAL APPLICATION

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Assistant Commissioner for Patents  
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Sir:

This is a request for filing a PROVISIONAL APPLICATION UNDER 37 CFR 1.53(b)(2).

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TITLE OF THE INVENTION			
Tumor Necrosis Factor Receptors 5, 6 Alpha and 6 Beta			

In connection with this application, the following are enclosed:

- 116 Pages of Specification (Optional: ☒ Abstract (Pg 123) ☒ Claims 22 (Pgs 117-122))
- 15 Sheets of Drawings
- Assignment to: \_\_\_\_\_
- Statement of Small Entity Status
- ☒ Other: Sequence Listings (pgs 81-116)/Diskette

The fee has been calculated as shown below. (Small entity fees indicated in parentheses.)

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☒ No ☐ Yes, the name of the U.S. Government agency and the Government contract number are:

The Commissioner is hereby authorized to charge any fees to Deposit Account No. 08-3425.

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## **Tumor Necrosis Factor Receptors 5, 6 $\alpha$ & 6 $\beta$**

### ***Field of the Invention***

5 The present invention relates to novel human genes encoding polypeptides which are members of the TNF receptor family. More specifically, isolated nucleic acid molecules are provided encoding human polypeptides named tumor necrosis factor receptor-5, -6 $\alpha$  & -6 $\beta$  hereinafter sometimes referred to as "TNFR-5, -6 $\alpha$ , & -6 $\beta$ " or generically as "TNFR polypeptides". TNFR polypeptides are also provided, as are vectors, host cells and recombinant methods for producing the same. The invention further relates to screening methods for identifying agonists and antagonists of TNFR polypeptide activity. Also provided are diagnostic and therapeutic methods utilizing such compositions.

### ***Background of the Invention***

15 Many biological actions, for instance, response to certain stimuli and natural biological processes, are controlled by factors, such as cytokines. Many cytokines act through receptors by engaging the receptor and producing an intracellular response.

20 For example, tumor necrosis factors (TNF) alpha and beta are cytokines which act through TNF receptors to regulate numerous biological processes, including protection against infection and induction of shock and inflammatory disease. The TNF molecules belong to the "TNF-ligand" superfamily, and act together with their receptors or counter-ligands, the "TNF-receptor" superfamily. So far, nine members of the TNF ligand superfamily have been identified and ten members of the TNF-receptor superfamily have been characterized.

25 Among the ligands there are included TNF- $\alpha$ , lymphotoxin- $\alpha$  (LT- $\alpha$ , also known as TNF- $\beta$ ), LT- $\beta$  (found in complex heterotrimer LT- $\alpha$ 2- $\beta$ ), FasL, CD40L, CD27L, CD30L, 4-1BBL, OX40L and nerve growth factor (NGF). The superfamily of TNF receptors includes the p55TNF receptor, p75TNF receptor, TNF receptor-related protein, FAS antigen or APO-1, CD40, CD27,

30



CD30, 4-1BB, OX40, low affinity p75 and NGF-receptor (Meager, A.,  
Biologicals, 22:291-295 (1994)).

Many members of the TNF-ligand superfamily are expressed by  
activated T-cells, implying that they are necessary for T-cell interactions with  
other cell types which underlie cell ontogeny and functions. (Meager, A.,  
supra).

Considerable insight into the essential functions of several members of  
the TNF receptor family has been gained from the identification and creation of  
mutants that abolish the expression of these proteins. For example, naturally  
occurring mutations in the FAS antigen and its ligand cause lymphoproliferative  
disease (Watanabe-Fukunaga, R., et al., Nature 356:314 (1992)), perhaps  
reflecting a failure of programmed cell death. Mutations of the CD40 ligand  
cause an X-linked immunodeficiency state characterized by high levels of  
immunoglobulin M and low levels of immunoglobulin G in plasma, indicating  
faulty T-cell-dependent B-cell activation (Allen, R.C. et al., Science 259:990  
(1993)). Targeted mutations of the low affinity nerve growth factor receptor  
cause a disorder characterized by faulty sensory innervation of peripheral  
structures (Lee, K.F. et al., Cell 69:737 (1992)).

TNF and LT- $\alpha$  are capable of binding to two TNF receptors (the 55-  
and 75-kd TNF receptors). A large number of biological effects elicited by  
TNF and LT- $\alpha$ , acting through their receptors, include hemorrhagic necrosis of  
transplanted tumors, cytotoxicity, a role in endotoxic shock, inflammation,  
immunoregulation, proliferation and anti-viral responses, as well as protection  
against the deleterious effects of ionizing radiation. TNF and LT- $\alpha$  are involved  
in the pathogenesis of a wide range of diseases, including endotoxic shock,  
cerebral malaria, tumors, autoimmune disease, AIDS and graft-host rejection  
(Beutler, B. and Von Huffel, C., Science 264:667-668 (1994)). Mutations in  
the p55 Receptor cause increased susceptibility to microbial infection.

Moreover, an about 80 amino acid domain near the C-terminus of  
TNFR1 (p55) and Fas was reported as the "death domain," which is  
responsible for transducing signals for programmed cell death (Tartaglia et al.,  
Cell 74:845 (1993)).

Apoptosis, or programmed cell death, is a physiologic process essential  
to the normal development and homeostasis of multicellular organisms (H.  
Steller, Science 267, 1445-1449 (1995)). Derangements of apoptosis  
contribute to the pathogenesis of several human diseases including cancer,  
neurodegenerative disorders and acquired immune deficiency syndrome (C.B.  
Thompson, Science 267, 1456-1462 (1995)). Recently, much attention has

focused on the signal transduction and biological function of two cell surface death receptors, Fas/APO-1 and TNFR-1 (J.L. Cleveland, J.N. Ihle, *Cell* 81, 479-482 (1995); A. Fraser, G. Evan, *Cell* 85, 781-784 (1996); S. Nagata, P. Golstein, *Science* 267, 1449-56 (1995)). Both are members of the TNF receptor family which also include TNFR-2, low affinity NGFR, CD40, and CD30, among others (C.A. Smith, et al., *Science* 248, 1019-23 (1990); M. Tewari, V.M. Dixit, in *Modular Texts in Molecular and Cell Biology* M. Purton, Heldin, Carl, Ed. (Chapman and Hall, London, 1995). While family members are defined by the presence of cysteine-rich repeats in their extracellular domains, Fas/APO-1 and TNFR-1 also share a region of intracellular homology, appropriately designated the "death domain", which is distantly related to the *Drosophila* suicide gene, reaper (P. Golstein, D. Marguet, V. Depraetere, *Cell* 81, 185-6 (1995); K. White et al., *Science* 264, 677-83 (1994)). This shared death domain suggests that both receptors interact with a related set of signal transducing molecules that, until recently, remained unidentified. Activation of Fas/APO-1 recruits the death domain-containing adapter molecule FADD/MORT1 (A.M. Chinnaiyan, K. O'Rourke, M. Tewari, V. M. Dixit, *Cell* 81, 505-12 (1995); M. P. Boldin, et al., *J. Biol Chem* 270, 7795-8 (1995); F.C. Kischkel, et al., *EMBO* 14, 5579-5588 (1995)), which in turn binds and presumably activates FLICE/MACH1, a member of the ICE/CED-3 family of pro-apoptotic proteases (M. Muzio et al., *Cell* 85, 817-827 (1996); M.P. Boldin, T.M. Goncharov, Y.V. Goltsev, D. Wallach, *Cell* 85, 803-815 (1996)). While the central role of Fas/APO-1 is to trigger cell death, TNFR-1 can signal an array of diverse biological activities-many of which stem from its ability to activate NF-kB (L.A. Tartaglia, D.V. Goeddel, *Immunol Today* 13, 151-3 (1992)). Accordingly, TNFR-1 recruits the multivalent adapter molecule TRADD, which like FADD, also contains a death domain (H. Hsu, J. Xiong, D.V. Goeddel, *Cell* 81, 495-504 (1995); H. Hsu, H.-B. Shu, M.-P. Pan, D.V. Goeddel, *Cell* 84, 299-308 (1996)). Through its associations with a number of signaling molecules including FADD, TRAF2, and RIP, TRADD can signal both apoptosis and NF-kB activation (H. Hsu, H.-B. Shu, M.-P. Pan, D.V. Goeddel, *Cell* 84, 299-308 (1996); H. Hsu, J. Huang, H.-B. Shu, V. Baichwal, D.V. Goeddel, *Immunity* 4, 387-396 (1996)).

The effects of TNF family ligands and TNF family receptors are varied and influence numerous functions, both normal and abnormal, in the biological processes of the mammalian system. There is a clear need, therefore, for identification and characterization of such receptors and ligands that influence biological activity, both normally and in disease states. In particular, there is a need to isolate and characterize novel members of the TNF receptor family.

### *Summary of the Invention*

The present invention provides isolated nucleic acid molecules comprising a polynucleotide encoding at least a portion of a TNFR-5, -6 $\alpha$  or -6 $\beta$  polypeptide having the complete amino acid sequences shown in SEQ ID NOS:2, 4 and 6, respectively, or the complete amino acid sequence encoded by a cDNA clone deposited in a bacterial host as ATCC Deposit Number 97798, 97810 and 97809, respectively. The nucleotide sequence determined by sequencing the deposited TNFR-5, -6 $\alpha$  and -6 $\beta$  clones, which are shown in Figures 1, 2 and 3 (SEQ ID NOS:1, 3 and 5, respectively), contain open reading frames encoding complete polypeptides of 259, 300 and 170 amino acid residues, respectively, including an initiation codon encoding an N-terminal methionine at nucleotide positions 183-185, 25-27, and 73-75 in SEQ ID NOS: 1, 3, and 5 respectively.

The TNFR proteins of the present invention share sequence homology with other TNF receptors. TNFR-5 shows the highest degree of sequence homology with the translation product for the human mRNA for nerve growth factor receptor (Figure 4) (SEQ ID NO:9), including multiple conserved cysteine rich domains. Splice variants TNFR-6 $\alpha$  and -6 $\beta$  show the highest degree of sequence homology with the translation products of the human mRNAs for TNFR-I and -II (Figure 4) (SEQ ID NOS:7 and 8, respectively) also including multiple conserved cysteine rich domains.

The TNFR-5, -6 $\alpha$  and -6 $\beta$  polypeptides have predicted leader sequences of 26, 30 and 30 amino acids, respectively; and the amino acid sequence of the predicted mature TNFR-5, -6 $\alpha$  and -6 $\beta$  polypeptides are also

shown in Figures 1, 2 and 3 as amino acid residues 27-259 (SEQ ID NO:2), 31-300 (SEQ ID NO:4), and 31-170 (SEQ ID NO:6), respectively.

Thus, one aspect of the invention provides an isolated nucleic acid molecule comprising a polynucleotide having a nucleotide sequence selected from the group consisting of: (a) a nucleotide sequence encoding a TNFR polypeptide having the complete amino acid sequence in SEQ ID NO:2, 4 or 6, or as encoded by the cDNA clone contained in ATCC Deposit No. 97798, 97810 or 97809; (b) a nucleotide sequence encoding a mature TNFR polypeptide having the amino acid sequence at positions 27-259 in SEQ ID NO:2, 31-300 in SEQ ID NO:4, or 31-170 in SEQ ID NO:6, or as encoded by the cDNA clone contained in ATCC Deposit No. 97798, 97810 or 97809; (c) a nucleotide sequence encoding a soluble extracellular domain of a TNFR polypeptide having the amino acid sequence at positions 27 to 240 in SEQ ID NO:2, 31 to 283 in SEQ ID NO:4, or 31 to 166 in SEQ ID NO:6, or as encoded by the cDNA clone contained in the ATCC Deposit No. 97798, 97810 or 97809; and (d) a nucleotide sequence complementary to any of the nucleotide sequences in (a), (b), or (c) above.

Further embodiments of the invention include isolated nucleic acid molecules that comprise a polynucleotide having a nucleotide sequence at least 90% identical, and more preferably at least 95%, 96%, 97%, 98% or 99% identical, to any of the nucleotide sequences in (a), (b), (c) and (d) above, or a polynucleotide which hybridizes under stringent hybridization conditions to a polynucleotide in (a), (b), (c) or (d), above. This polynucleotide which hybridizes does not hybridize under stringent hybridization conditions to a polynucleotide having a nucleotide sequence consisting of only A residues or of only T residues. An additional nucleic acid embodiment of the invention relates to an isolated nucleic acid molecule comprising a polynucleotide which encodes the amino acid sequence of an epitope-bearing portion of a TNFR polypeptide having an amino acid sequence in (a), (b) or (c), above.

The present invention also relates to recombinant vectors, which include the isolated nucleic acid molecules of the present invention, and to host cells containing the recombinant vectors, as well as to methods of making such vectors and host cells and for using them for production of TNFR polypeptides or peptides by recombinant techniques.

The invention further provides an isolated TNFR polypeptide comprising an amino acid sequence selected from the group consisting of: (a) the amino acid sequence of a full-length TNFR polypeptide having the complete amino acid sequence shown in SEQ ID NO:2, 4 or 6 or as encoded by the cDNA clone contained in ATCC Deposit No. 97798, 97810 or 97809; (b) the amino acid sequence of a mature TNFR polypeptide having the amino acid sequence at positions 27-259 in SEQ ID NO:2, 31-300 in SEQ ID NO:4, or 31-170 in SEQ ID NO:6, or as encoded by the cDNA clone contained in ATCC Deposit No. 97798, 97810 or 97809; or (c) the amino acid sequence of a soluble extracellular domain of a TNFR polypeptide having the amino acid sequence at positions 27 to 240 in SEQ ID NO:2, 31 to 283 in SEQ ID NO:4, or 31 to 166 in SEQ ID NO:6, or as encoded by the cDNA clone contained in ATCC Deposit No. 97798, 97810 or 97809.

The polypeptides of the present invention also include polypeptides having an amino acid sequence at least 80% identical, more preferably at least 90% identical, and still more preferably 95%, 96%, 97%, 98% or 99% identical to those described in (a), (b) or (c) above, as well as polypeptides having an amino acid sequence with at least 90% similarity, and more preferably at least 95% similarity, to those above.

An additional embodiment of this aspect of the invention relates to a peptide or polypeptide which comprises the amino acid sequence of an epitope-bearing portion of a TNFR polypeptide having an amino acid sequence described in (a), (b) or (c), above. Peptides or polypeptides having the amino acid sequence of an epitope-bearing portion of a TNFR polypeptide of the invention include portions of such polypeptides with at least six or

seven, preferably at least nine, and more preferably at least about 30 amino acids to about 50 amino acids, although epitope-bearing polypeptides of any length up to and including the entire amino acid sequence of a polypeptide of the invention described above also are included in the invention.

5 In another embodiment, the invention provides an isolated antibody that binds specifically to a TNFR polypeptide having an amino acid sequence described in (a), (b) or (c) above. The invention further provides methods for isolating antibodies that bind specifically to a TNFR polypeptide having an amino acid sequence as described herein. Such antibodies are useful  
10 diagnostically or therapeutically as described below.

Tumor Necrosis Factor (TNF) family ligands are known to be among the most pleiotropic cytokines, inducing a large number of cellular responses, including cytotoxicity, anti-viral activity, immunoregulatory activities, and the transcriptional regulation of several genes. The invention also provides for  
15 pharmaceutical compositions comprising TNFR polypeptides, particularly human TNFR polypeptides, which may be employed, for instance, to treat infectious disease including HIV infection, endotoxic shock, cancer, autoimmune diseases, graft vs. host disease, acute graft rejection, chronic graft rejection, neurodegenerative disorders, myelodysplastic syndromes, ischemic  
20 injury, toxin-induced liver disease, septic shock, cachexia and anorexia. Methods of treating individuals in need of TNFR polypeptides are also provided.

The invention further provides compositions comprising a TNFR polynucleotide or a TNFR polypeptide for administration to cells *in vitro*, to  
25 cells *ex vivo* and to cells *in vivo*, or to a multicellular organism. In certain particularly preferred embodiments of this aspect of the invention, the compositions comprise a TNFR polynucleotide for expression of a TNFR polypeptide in a host organism for treatment of disease. Particularly preferred in this regard is expression in a human patient for treatment of a dysfunction.  
30 associated with aberrant endogenous activity of a TNFR polypeptide.

In another aspect, a screening assay for agonists and antagonists is provided which involves determining the effect a candidate compound has on TNFR polypeptide binding to a TNF-family ligand. In particular, the method involves contacting the TNF-family ligand with a TNFR polypeptide and a candidate compound and determining whether TNFR polypeptide binding to the TNF-family ligand is increased or decreased due to the presence of the candidate compound. In this assay, an increase in binding of a TNFR polypeptide over the standard binding indicates that the candidate compound is an agonist of TNFR polypeptide binding activity and a decrease in TNFR polypeptide binding compared to the standard indicates that the compound is an antagonist of TNFR polypeptide binding activity.

It has been discovered that TNFR-5 is expressed not only in prostate tissue but also in endothelial cells, stimulated monocytes and keratinocytes. TNFR-6 $\alpha$  and -6 $\beta$  are expressed endothelial cells, keratinocytes, normal prostate and prostate tumor tissue. For a number of disorders of these tissues or cell, particularly of the immune system significantly higher or lower levels of TNFR gene expression may be detected in certain tissues (e.g., cancerous tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) taken from an individual having such a disorder, relative to a "standard" TNFR gene expression level, i.e., the TNFR expression level in healthy tissue from an individual not having the immune system disorder. Thus, the invention provides a diagnostic method useful during diagnosis of such a disorder, which involves: (a) assaying TNFR gene expression level in cells or body fluid of an individual; (b) comparing the TNFR gene expression level with a standard TNFR gene expression level, whereby an increase or decrease in the assayed TNFR gene expression level compared to the standard expression level is indicative of disorder in the immune system.

An additional aspect of the invention is related to a method for treating an individual in need of an increased level of TNFR polypeptide activity in the body comprising administering to such an individual a composition comprising

a therapeutically effective amount of an isolated TNFR polypeptide of the invention or an agonist thereof.

A still further aspect of the invention is related to a method for treating an individual in need of a decreased level of TNFR polypeptide activity in the body comprising, administering to such an individual a composition comprising a therapeutically effective amount of a TNFR antagonist. Preferred antagonists for use in the present invention are TNFR-specific antibodies.

### *Brief Description of the Figures*

Figure 1 shows the nucleotide sequence (SEQ ID NO:1) and deduced amino acid sequence (SEQ ID NO:2) of TNFR-5.

Figure 2 shows the nucleotide sequence (SEQ ID NO:3) and deduced amino acid sequence (SEQ ID NO:4) of TNFR-6 $\alpha$ .

Figure 3 shows the nucleotide sequence (SEQ ID NO:5) and deduced amino acid sequence (SEQ ID NO:6) of TNFR-6 $\beta$ .

Figure 4 shows an alignment created by the Clustal method using the Megaline program in the DNASTAR suite comparing the amino acid sequences of TNFR-5 ("TNFR-like"), TNFR-6 $\alpha$  ("TNFR-6a"), and TNFR-6 $\beta$  ("TNFR-6b") with other TNF receptors, as follows: TNFR1 (SEQ ID NO:7); TNFR2 (SEQ ID NO:8); NGFR (SEQ ID NO:9); LT $\beta$ R (SEQ ID NO:10); FAS (SEQ ID NO:11); CD27 (SEQ ID NO:12); CD30 (SEQ ID NO:13); CD40 (SEQ ID NO:14); 4-1BB (SEQ ID NO:15); OX40 (SEQ ID NO:16); VC22 (SEQ ID NO:17); and CRMB (SEQ ID NO:18).

Figures 5, 6 and 7 show separate analyses of the TNFR-5, 6 $\alpha$  and -6 $\beta$  amino acid sequences, respectively. Alpha, beta, turn and coil regions; hydrophilicity and hydrophobicity; amphipathic regions; flexible regions; antigenic index and surface probability are shown. In the "Antigenic Index - Jameson-Wolf" graphs, the indicate location of the highly antigenic regions of



the proteins, i.e., regions from which epitope-bearing fragments may be obtained.

Figure 8 shows the nucleotide sequence of fragments related to the TNFR genes of the present invention, including: HPRCB54R (SEQ ID NO:19), HSJAU57RA (SEQ ID NO:20), HELBP70R (SEQ ID NO:21), and HUSCB54R (SEQ ID NO:22) all of which are related to SEQ ID NO:1; and HELDI06R (SEQ ID NO:23) and HCEOW38R (SEQ ID NO:24) both of which are related to SEQ ID NOS:3 and 5.

### *Detailed Description*

The present invention provides isolated nucleic acid molecules comprising a polynucleotide encoding a TNFR-5, -6 $\alpha$  or -6 $\beta$  polypeptide, generically "TNFR polypeptide(s)" having the amino acid sequence shown in SEQ ID NOS:2, 4 and 6, respectively, which were determined by sequencing cloned cDNAs. The nucleotide sequences shown in Figures 1, 2 and 3 (SEQ ID NOS:1, 3 and 5) were obtained by sequencing the HPRCB54, HPHA52 and HTPCH84 clones, which were deposited on November 20, 1996, November 22, 1996, and November 22, 1996, respectively, at the American Type Culture Collection, 12301 Park Lawn Drive, Rockville, Maryland 20852, and given accession numbers ATCC 97798, 97810 and 97809, respectively. The deposited clones are contained in the pBluescript SK(-) plasmid (Stratagene, La Jolla, CA).

The TNFR-5 protein of the present invention has an amino acid sequence which is 21.7% identical to and shares multiple conserved cysteine rich domains with the translation product of the human nerve growth factor (hNGF) mRNA (SEQ ID NO:9) as illustrated in Figure 4. hNGF is thought to play an important role in the development, survival, apoptosis and function of neurons (Lee, F.K. et al., Cell 69:737) and lymphocytes (Torcia, M. et al., Cell 85:3369 (1996)).

The TNFR-6 $\alpha$  and -6 $\beta$  proteins of the present invention are splice variants which share an identical nucleotide and amino acid sequence over the N-terminal 142 residues of the respective proteins. The amino acid sequences of these proteins are about 23% similar to and share multiple conserved cysteine rich domains with the translation product of the human TNFR-2 mRNA (Figure 4) (SEQ ID NO:8). Importantly, these proteins share substantial sequence similarity over their extracellular domains including four repeated cysteine rich motifs with significant intersubunit homology. TNFR-2 is thought to exclusively mediate human T-cell proliferation by TNF (PCT WO/94/09137).

### *Nucleic Acid Molecules*

Unless otherwise indicated, all nucleotide sequences determined by sequencing a DNA molecule herein were determined using an automated DNA sequencer (such as the Model 373 from Applied Biosystems, Inc., Foster City, CA), and all amino acid sequences of polypeptides encoded by DNA molecules determined herein were predicted by translation of a DNA sequence determined as above. Therefore, as is known in the art for any DNA sequence determined by this automated approach, any nucleotide sequence determined herein may contain some errors. Nucleotide sequences determined by automation are typically at least about 90% identical, more typically at least about 95% to at least about 99.9% identical to the actual nucleotide sequence of the sequenced DNA molecule. The actual sequence can be more precisely determined by other approaches including manual DNA sequencing methods well known in the art. As is also known in the art, a single insertion or deletion in a determined nucleotide sequence compared to the actual sequence will cause a frame shift in translation of the nucleotide sequence such that the predicted amino acid sequence encoded by a determined nucleotide sequence will be completely different from the amino acid sequence actually encoded by

the sequenced DNA molecule, beginning at the point of such an insertion or deletion.

By "nucleotide sequence" of a nucleic acid molecule or polynucleotide is intended, for a DNA molecule or polynucleotide, a sequence of deoxyribonucleotides, and for an RNA molecule or polynucleotide, the corresponding sequence of ribonucleotides (A, G, C and U), where each thymidine deoxyribonucleotide (T) in the specified deoxyribonucleotide sequence is replaced by the ribonucleotide uridine (U).

Using the information provided herein, such as the nucleotide sequences in Figures 1, 2 and 3 (SEQ ID NOS:1, 3 and 5), a nucleic acid molecule of the present invention encoding a TNFR polypeptide may be obtained using standard cloning and screening procedures, such as those for cloning cDNAs using mRNA as starting material. Illustrative of the invention, the TNFR-5 nucleic acid molecule described in Figure 1 was discovered in a cDNA library derived from prostate tissue. Additional clones of the same gene were also identified in cDNA libraries from the following tissues: endothelial cells, stimulated monocytes, and keratinocytes. TNFR-6 $\alpha$  and -6 $\beta$  clones (Figures 2 and 3, respectively) were identified in cDNA libraries from the following tissues: endothelial cells, keratinocytes, normal prostate tissue, and prostate tumor tissue.

The determined nucleotide sequences of the TNFR cDNAs of Figures 1, 2 and 3 (SEQ ID NOS:1, 3 and 5) contain open reading frames encoding proteins of 259, 300 and 170 amino acid residues, with an initiation codon at nucleotide positions 183-185, 25-27, 73-75 of the nucleotide sequences in Figures 1, 2 and 3 (SEQ ID NOS:1, 3 and 5), respectively.

The open reading frames of the TNFR-6 $\alpha$  and -6 $\beta$  genes share sequence homology with the translation product of the human mRNA for TNFR-2, including the soluble extracellular domain of about residues 31-283 of SEQ ID NO:4 and 31-166 of SEQ ID NO:6, respectively. The open reading frame of the TNFR-5 gene shares sequence homology with the translation product of the human mRNA for NGFR, including the following conserved

domains: (a) a soluble extracellular domain of about 214 amino acids (residues 27-240 of SEQ ID NO:2); and (b) a transmembrane domain of about 19 amino acids (residues 241-259 of SEQ ID NO:2).

As one of ordinary skill would appreciate, due to the possibilities of sequencing errors discussed above, the actual complete TNFR polypeptides encoded by the deposited cDNAs, which comprise about 259, 300, and 170 amino acids, may be somewhat longer or shorter. More generally, the actual open reading frames may be anywhere in the range of  $\pm 20$  amino acids, more likely in the range of  $\pm 10$  amino acids, of that predicted from the first methionine codon from the N-terminus shown in Figures 1, 2 and 3 (SEQ ID NOS:1, 3 and 5), which is in-frame with the translated sequences shown in each respective figure. It will further be appreciated that, depending on the analytical criteria used for identifying various functional domains, the exact "address" of the extracellular and transmembrane domain(s) of the TNFR polypeptides may differ slightly from the predicted positions above. For example, the exact location of the extracellular domain in SEQ ID NO:2 may vary slightly (e.g., the address may "shift" by about 1 to about 20 residues, more likely about 1 to about 5 residues) depending on the criteria used to define the domain. In this case, the beginning of the transmembrane domain and the end of the extracellular domain were predicted on the basis of the identification of the hydrophobic amino acid sequence in the above indicated positions, as shown in Figure 5. In any event, as discussed further below, the invention further provides polypeptides having various residues deleted from the N-terminus of the complete polypeptide, including polypeptides lacking one or more amino acids from the N-terminus of the extracellular domain described herein, which constitute soluble forms of the extracellular domains of the TNFR-5, -6 $\alpha$  & -6 $\beta$  proteins.

### *Leader and Mature Sequences*

The amino acid sequences of the complete TNFR proteins include a leader sequence and a mature protein, as shown in SEQ ID NOS:2, 4 and 6. More in particular, the present invention provides nucleic acid molecules encoding mature forms of the TNFR proteins. Thus, according to the signal hypothesis, once export of the growing protein chain across the rough endoplasmic reticulum has been initiated, proteins secreted by mammalian cells have a signal or secretory leader sequence which is cleaved from the complete polypeptide to produce a secreted "mature" form of the protein. Most mammalian cells and even insect cells cleave secreted proteins with the same specificity. However, in some cases, cleavage of a secreted protein is not entirely uniform, which results in two or more mature species of the protein. Further, it has long been known that the cleavage specificity of a secreted protein is ultimately determined by the primary structure of the complete protein, that is, it is inherent in the amino acid sequence of the polypeptide. Therefore, the present invention provides a nucleotide sequence encoding a mature TNFR polypeptide having the amino acid sequence encoded by a cDNA clone contained in a host identified as ATCC Deposit No. 97798, 97810 or 97809. By the "mature TNFR polypeptides having the amino acid sequence encoded by a cDNA clone in ATCC Deposit No. 97798, 97810, or 97809" is meant the mature form(s) of the protein produced by expression in a mammalian cell (e.g., COS cells, as described below) of the complete open reading frame encoded by the human DNA sequence of the clone contained in the vector in the deposited host.

In addition, methods for predicting whether a protein has a secretory leader as well as the cleavage point for that leader sequence are available. For instance, the method of McGeoch (*Virus Res.* 3:271-286 (1985)) uses the information from a short N-terminal charged region and a subsequent uncharged region of the complete (uncleaved) protein. The method of von Heinje (*Nucleic Acids Res.* 14:4683-4690 (1986)) uses the information from the

residues surrounding the cleavage site, typically residues -13 to +2 where +1 indicates the amino terminus of the mature protein. The accuracy of predicting the cleavage points of known mammalian secretory proteins for each of these methods is in the range of 75-80% (von Heinje, *supra*). However, the two methods do not always produce the same predicted cleavage point(s) for a given protein.

In the present case, the deduced amino acid sequence of the complete TNFR polypeptides were analyzed by a computer program "PSORT", available from Dr. Kenta . . . of the Institute for Chemical Research, Kyoto University (see K. Nakai and M. Kanehisa, *Genomics* 14:897-911 (1992)), which is an expert system for predicting the cellular location of a protein based on the amino acid sequence. As part of this computational prediction of localization, the methods of McGeoch and von Heinje are incorporated. The analysis of the TNFR amino acid sequences by this program provided the following results: TNFR-5, -6 $\alpha$  & -6 $\beta$  encode mature polypeptides having the amino acid sequences of residues 27-259, 31-300 and 31-170 of SEQ ID NOS:2, 4 and 6, respectively.

As indicated, nucleic acid molecules of the present invention may be in the form of RNA, such as mRNA, or in the form of DNA, including, for instance, cDNA and genomic DNA obtained by cloning or produced synthetically. The DNA may be double-stranded or single-stranded. Single-stranded DNA or RNA may be the coding strand, also known as the sense strand, or it may be the non-coding strand, also referred to as the anti-sense strand.

By "isolated" nucleic acid molecule(s) is intended a nucleic acid molecule, DNA or RNA, which has been removed from its native environment. For example, recombinant DNA molecules contained in a vector are considered isolated for the purposes of the present invention. Further examples of isolated DNA molecules include recombinant DNA molecules maintained in heterologous host cells or purified (partially or substantially) DNA molecules

in solution. Isolated RNA molecules include *in vivo* or *in vitro* RNA transcripts of the DNA molecules of the present invention. Isolated nucleic acid molecules according to the present invention further include such molecules produced synthetically.

5 Isolated nucleic acid molecules of the present invention include DNA molecules comprising an open reading frame (ORF) with an initiation codon at positions 183-185, 25-27 73-75 of the nucleotide sequences shown in SEQ ID NOS:1, 3 and 5 respectively.

10 Also included are DNA molecules comprising the coding sequence for the predicted mature TNFR polypeptides shown at positions 27-259, 31-300, and 31-170 of SEQ ID NOS:2, 4 and 6, respectively.

15 In addition, isolated nucleic acid molecules of the invention include DNA molecules which comprise a sequence substantially different from those described above but which, due to the degeneracy of the genetic code, still encode a TNFR protein. Of course, the genetic code and species-specific codon preferences are well known in the art. Thus, it would be routine for one skilled in the art to generate the degenerate variants described above. for instance, to optimize codon expression for a particular host (e.g., change codons in the human mRNA to those preferred by a bacterial host such as *E. coli*).

20 In another aspect, the invention provides isolated nucleic acid molecules encoding a TNFR polypeptide having an amino acid sequence encoded by the cDNA clone contained in the plasmid deposited as ATCC Deposit No. 97798, 97810, or 97809. Preferably, this nucleic acid molecule will encode the mature polypeptide encoded by the above-described deposited cDNA clone.

25 The invention further provides an isolated nucleic acid molecule having the nucleotide sequence shown in Figure 1, 2 or 3 (SEQ ID NO:1, 3 or 5) or the nucleotide sequence of the TNFR cDNAs contained in the above-described

deposited clones, or a nucleic acid molecule having a sequence complementary to one of the above sequences. Such isolated molecules, particularly DNA molecules, are useful as probes for gene mapping, by *in situ* hybridization with chromosomes, and for detecting expression of the TNFR genes in human tissue, for instance, by Northern blot analysis.

The present invention is further directed to nucleic acid molecules encoding portions of the nucleotide sequences described herein as well as to fragments of the isolated nucleic acid molecules described herein. In particular, the invention provides polynucleotides having a nucleotide sequence representing the portion of SEQ ID NO:1, 3 or 5 which consist of positions 183-959, 25-924 and 73-582 of SEQ ID NOS:1, 3 and 5, respectively. Also contemplated are polynucleotides encoding TNFR polypeptides which lack an amino terminal methionine such polynucleotides having a nucleotide sequence representing the portion of SEQ ID NOS:1, 3 and 5 which consist of positions 186-959, 28-924, and 76-582 of SEQ ID NOS:1,3 and 5, respectively. Polypeptides encoded by such polynucleotides are also provided, such polypeptides comprising an amino acid sequence at positions 2-259, 2-300, and 2-170 of SEQ ID NOS:2, 4, and 6, respectively.

In addition, the invention provides nucleic acid molecules having nucleotide sequences related to extensive portions of SEQ ID NOS:1, 3 and 5 which have been determined from the following related cDNA clones: HELBP70R (SEQ ID NO:21), HPRCB54R (SEQ ID NO:19), HSJAU57RA (SEQ ID NO:20) and HUSCB54R (SEQ ID NO:22) are related to SEQ ID NO:1; HELDI06R (SEQ ID NO:23) and HCEOW38R (SEQ ID NO:24) are related to SEQ ID NOS:3 and 5. The nucleotide sequences of each of these clones is shown in Figure 8.

More generally, by a fragment of an isolated nucleic acid molecule having the nucleotide sequence of the deposited cDNA or the nucleotide sequence shown in Figures 1, 2 or 3 (SEQ ID NOS:1, 3 or 5) is intended fragments at least about 15 nt, and more preferably at least about 20 nt, still



more preferably at least about 30 nt, and even more preferably, at least about 40 nt in length which are useful as diagnostic probes and primers as discussed herein. Of course, larger fragments 50-300 nt in length are also useful according to the present invention as are fragments corresponding to most, if not all, of the nucleotide sequence of the deposited cDNAs or as shown in Figures 1, 2 and 3 (SEQ ID NOS:1, 3 and 5). By a fragment at least 20 nt in length, for example, is intended fragments which include 20 or more contiguous bases from the nucleotide sequence of a deposited cDNA or the nucleotide sequence as shown in Figures 1, 2 and 3 (SEQ ID NOS:1, 3 and 5). Preferred nucleic acid fragments of the present invention include nucleic acid molecules encoding epitope-bearing portions of the TNFR polypeptides as identified in Figures 5, 6 and 7 and described in more detail below.

In another aspect, the invention provides an isolated nucleic acid molecule comprising a polynucleotide which hybridizes under stringent hybridization conditions to a portion of the polynucleotide in a nucleic acid molecule of the invention described above, for instance, a cDNA clone contained in ATCC Deposit No. 97798, 97810 or 97809. By "stringent hybridization conditions" is intended overnight incubation at 42° C in a solution comprising: 50% formamide, 5x SSC (150 mM NaCl, 15 mM trisodium citrate), 50 mM sodium phosphate (pH 7.6), 5x Denhardt's solution, 10% dextran sulfate, and 20 µg/ml denatured, sheared salmon sperm DNA, followed by washing the filters in 0.1x SSC at about 65° C.

By a polynucleotide which hybridizes to a "portion" of a polynucleotide is intended a polynucleotide (either DNA or RNA) hybridizing to at least about 15 nucleotides (nt), and more preferably at least about 20 nt, still more preferably at least about 30 nt, and even more preferably about 30-70 (e.g., 50) nt of the reference polynucleotide. These are useful as diagnostic probes and primers as discussed above and in more detail below.

By a portion of a polynucleotide of "at least 20 nt in length," for example, is intended 20 or more contiguous nucleotides from the nucleotide

sequence of the reference polynucleotide (e.g., a deposited cDNA or a nucleotide sequence as shown in Figure 1, 2 or 3 (SEQ ID NO:1, 3 or 5)). Of course, a polynucleotide which hybridizes only to a poly A sequence (such as the 3' terminal poly(A) tract of a TNFR cDNA, or to a complementary stretch of T (or U) residues, would not be included in a polynucleotide of the invention used to hybridize to a portion of a nucleic acid of the invention, since such a polynucleotide would hybridize to any nucleic acid molecule containing a poly (A) stretch or the complement thereof (e.g., practically any double-stranded cDNA clone).

As indicated, nucleic acid molecules of the present invention which encode a TNFR polypeptide may include, but are not limited to those encoding the amino acid sequence of the mature polypeptide, by itself; and the coding sequence for the mature polypeptide and additional sequences, such as those encoding the about 26-31 amino acid leader or secretory sequence, such as a pre-, or pro- or prepro- protein sequence; the coding sequence of the mature polypeptide, with or without the aforementioned additional coding sequences.

Also encoded by nucleic acids of the invention are the above protein sequences together with additional, non-coding sequences, including for example, but not limited to introns and non-coding 5' and 3' sequences, such as the transcribed, non-translated sequences that play a role in transcription, mRNA processing, including splicing and polyadenylation signals, for example - ribosome binding and stability of mRNA; an additional coding sequence which codes for additional amino acids, such as those which provide additional functionalities.

Thus, the sequence encoding the polypeptide may be fused to a marker sequence, such as a sequence encoding a peptide which facilitates purification of the fused polypeptide. In certain preferred embodiments of this aspect of the invention, the marker amino acid sequence is a hexa-histidine peptide, such as the tag provided in a pQE vector (QIAGEN, Inc., 9259 Eton Avenue,

Chatsworth, CA, 91311), among others, many of which are commercially available. As described in Gentz *et al.*, *Proc. Natl. Acad. Sci. USA* 86:821-824 (1989), for instance, hexa-histidine provides for convenient purification of the fusion protein. The "HA" tag is another peptide useful for purification which corresponds to an epitope derived from the influenza hemagglutinin protein, which has been described by Wilson *et al.*, *Cell* 37: 767 (1984). As discussed below, other such fusion proteins include a TNFR-5, -6 $\alpha$  or -6 $\beta$  fused to Fc at the N- or C-terminus.

#### ***Variant and Mutant Polynucleotides***

The present invention further relates to variants of the nucleic acid molecules of the present invention, which encode portions, analogs or derivatives of a TNFR polypeptide. Variants may occur naturally, such as a natural allelic variant. By an "allelic variant" is intended one of several alternate forms of a gene occupying a given locus on a chromosome of an organism. *Genes II*, Lewin, B., ed., John Wiley & Sons, New York (1985). Non-naturally occurring variants may be produced using art-known mutagenesis techniques.

Such variants include those produced by nucleotide substitutions, deletions or additions. The substitutions, deletions or additions may involve one or more nucleotides. The variants may be altered in coding regions, non-coding regions, or both. Alterations in the coding regions may produce conservative or non-conservative amino acid substitutions, deletions or additions. Especially preferred among these are silent substitutions, additions and deletions, which do not alter the properties and activities of the TNFR polypeptide or portions thereof. Also especially preferred in this regard are conservative substitutions.

Highly preferred are nucleic acid molecules encoding a mature protein having an amino acid sequence shown in SEQ ID NOS:2, 4 and 6 or the mature TNFR polypeptide sequences encoded by the deposited cDNA clones.

Most highly preferred are nucleic acid molecules encoding the extracellular domain of a protein having the amino acid sequence shown in SEQ ID NO:2, 4 or 6 or the extracellular domain of a TNFR amino acid sequence encoded by a deposited cDNA clone.

5 Further embodiments include an isolated nucleic acid molecule comprising a polynucleotide having a nucleotide sequence at least 90% identical, and more preferably at least 95%, 96%, 97%, 98% or 99% identical to a polynucleotide selected from the group consisting of: (a) a nucleotide sequence encoding a TNFR polypeptide having the complete amino acid  
10 sequence in SEQ ID NO:2, 4 or 6, or as encoded by a cDNA clone contained in ATCC Deposit No. 97798, 97810, or 97809; (b) a nucleotide sequence encoding a mature TNFR polypeptide having an amino acid sequence at positions 27-259, 31-300 or 31-170 in SEQ ID NO:2, 4 or 6, respectively, or as encoded by a cDNA clone contained in ATCC Deposit No. 97798, 97810  
15 or 97809; (c) a nucleotide sequence encoding a soluble extracellular domain of a TNFR polypeptide having the amino acid sequence at positions 27-240, 31-283, and 31-166 of SEQ ID NOS:2, 4 and 6, respectively; and (d) a nucleotide sequence complementary to any of the nucleotide sequences in (a), (b) or (c) above.

20 Further embodiments of the invention include isolated nucleic acid molecules that comprise a polynucleotide having a nucleotide sequence at least 90% identical, and more preferably at least 95%, 96%, 97%, 98% or 99% identical, to any of the nucleotide sequences in (a), (b), (c), or (d), above, or a polynucleotide which hybridizes under stringent hybridization conditions to a  
25 polynucleotide in (a), (b), (c), or (d), above. This polynucleotide which hybridizes does not hybridize under stringent hybridization conditions to a polynucleotide having a nucleotide sequence consisting of only A residues or of only T residues. An additional nucleic acid embodiment of the invention relates to an isolated nucleic acid molecule comprising a polynucleotide which

encodes the amino acid sequence of an epitope-bearing portion of a TNFR polypeptide having an amino acid sequence in (a), (b), (c), or (d), above.

The present invention also relates to recombinant vectors, which include the isolated nucleic acid molecules of the present invention, and to host  
5 cells containing the recombinant vectors, as well as to methods of making such vectors and host cells and for using them for production of TNFR polypeptides or peptides by recombinant techniques.

By a polynucleotide having a nucleotide sequence at least, for example, 95% "identical" to a reference nucleotide sequence encoding a TNFR  
10 polypeptide is intended that the nucleotide sequence of the polynucleotide is identical to the reference sequence except that the polynucleotide sequence may include up to five point mutations per each 100 nucleotides of the reference nucleotide sequence encoding the TNFR polypeptide. In other words, to obtain a polynucleotide having a nucleotide sequence at least 95%  
15 identical to a reference nucleotide sequence, up to 5% of the nucleotides in the reference sequence may be deleted or substituted with another nucleotide, or a number of nucleotides up to 5% of the total nucleotides in the reference sequence may be inserted into the reference sequence. These mutations of the reference sequence may occur at the 5' or 3' terminal positions of the reference  
20 nucleotide sequence or anywhere between those terminal positions, interspersed either individually among nucleotides in the reference sequence or in one or more contiguous groups within the reference sequence.

As a practical matter, whether any particular nucleic acid molecule is at least 90%, 95%, 96%, 97%, 98% or 99% identical to, for instance, a nucleotide  
25 sequence shown in Figure 1, 2 or 3, or to the nucleotides sequence of a deposited cDNA clone can be determined conventionally using known computer programs such as the Bestfit program (Wisconsin Sequence Analysis Package, Version 8 for Unix, Genetics Computer Group, University Research Park, 575 Science Drive, Madison, WI 53711). Bestfit uses the local  
30 homology algorithm of Smith and Waterman, *Advances in Applied Mathematics*

2:482-489 (1981), to find the best segment of homology between two sequences. When using Bestfit or any other sequence alignment program to determine whether a particular sequence is, for instance, 95% identical to a reference sequence according to the present invention, the parameters are set, of course, such that the percentage of identity is calculated over the full length of the reference nucleotide sequence and that gaps in homology of up to 5% of the total number of nucleotides in the reference sequence are allowed.

The present application is directed to nucleic acid molecules at least 90%, 95%, 96%, 97%, 98% or 99% identical to a nucleic acid sequence shown in Figure 1, 2 or 3 (SEQ ID NO:1, 3 or 5) or to the nucleic acid sequence of a deposited cDNA, irrespective of whether they encode a polypeptide having TNFR activity. This is because even where a particular nucleic acid molecule does not encode a polypeptide having TNFR activity, one of skill in the art would still know how to use the nucleic acid molecule, for instance, as a hybridization probe or a polymerase chain reaction (PCR) primer. Uses of the nucleic acid molecules of the present invention that do not encode a polypeptide having TNFR activity include, inter alia, (1) isolating a TNFR gene or allelic variants thereof in a cDNA library; (2) *in situ* hybridization (e.g., "FISH") to metaphase chromosomal spreads to provide precise chromosomal location of the TNFR gene, as described in Verma et al., Human Chromosomes: A Manual of Basic Techniques, Pergamon Press, New York (1988); and Northern Blot analysis for detecting TNFR mRNA expression in specific tissues.

Preferred, however, are nucleic acid molecules having sequences at least 90%, 95%, 96%, 97%, 98% or 99% identical to a nucleic acid sequence shown in Figure 1, 2 or 3 (SEQ ID NO:1, 3 or 5) or to the nucleic acid sequence of a deposited cDNA which do, in fact, encode polypeptides having TNFR protein activity. By "a polypeptide having TNFR activity" is intended polypeptides exhibiting activity similar, but not necessarily identical, to an activity of a mature or extracellular forms of a TNFR-5, -6 $\alpha$  or -6 $\beta$  protein of the

invention, as measured in a particular biological assay. The TNF family ligands induce various cellular responses by binding to TNF-family receptors, including the TNFR-5, -6 $\alpha$  and -6 $\beta$  of the present invention. Cells which express the TNFR proteins are believed to have a potent cellular response to TNFR-I receptor ligands including B lymphocytes (CD19+), both CD4 and CD8+ T lymphocytes, monocytes and endothelial cells. By a "cellular response to a TNF-family ligand" is intended any genotypic, phenotypic, and/or morphological change to a cell, cell line, tissue, tissue culture or patient that is induced by a TNF-family ligand. As indicated, such cellular responses include not only normal physiological responses to TNF-family ligands, but also diseases associated with increased cell proliferation or the inhibition of increased cell proliferation, such as by the inhibition of apoptosis.

Screening assays for the forgoing are known in the art. One such screening assay involves the use of cells which express the receptor (for example, transfected CHO cells) in a system which measures extracellular pH changes caused by receptor activation, for example, as described in Science 246:181-296 (October 1989). For example, a TNF-family ligand may be contacted with a cell which expresses the mature form of the receptor polypeptide of the present invention and a second messenger response, e.g., signal transduction or pH changes, may be measured to determine whether the TNFR polypeptide is active.

Of course, due to the degeneracy of the genetic code, one of ordinary skill in the art will immediately recognize that a large number of the nucleic acid molecules having a sequence at least 90%, 95%, 96%, 97%, 98%, or 99% identical to the nucleic acid sequence of a deposited cDNA or the nucleic acid sequence shown in Figure 1, 2 or 3 (SEQ ID NO:1, 3 or 5) will encode a polypeptide "having TNFR protein activity." In fact, since degenerate variants of these nucleotide sequences all encode the same polypeptide, this will be clear to the skilled artisan even without performing the above described comparison assay. It will be further recognized in the art that, for such nucleic acid molecules that are not degenerate variants, a reasonable number will also encode a polypeptide having TNFR protein activity. This is because the skilled artisan is fully aware of amino acid substitutions that are either less likely or not likely to significantly effect protein function (e.g., replacing one

aliphatic amino acid with a second aliphatic amino acid), as further described below.

#### ***Vectors and Host Cells***

The present invention also relates to vectors which include the isolated DNA molecules of the present invention, host cells which are genetically engineered with the recombinant vectors, and the production of TNFR polypeptides or fragments thereof by recombinant techniques. The vector may be, for example, a phage, plasmid, viral or retroviral vector. Retroviral vectors may be replication competent or replication defective. In the latter case, viral propagation generally will occur only in complementing host cells.

The polynucleotides may be joined to a vector containing a selectable marker for propagation in a host. Generally, a plasmid vector is introduced in a precipitate, such as a calcium phosphate precipitate, or in a complex with a charged lipid. If the vector is a virus, it may be packaged in vitro using an appropriate packaging cell line and then transduced into host cells.

The DNA insert should be operatively linked to an appropriate promoter, such as the phage lambda PL promoter, the *E. coli lac*, *trp*, *phoA* and *tac* promoters, the SV40 early and late promoters and promoters of retroviral LTRs, to name a few. Other suitable promoters will be known to the skilled artisan. The expression constructs will further contain sites for transcription initiation, termination and, in the transcribed region, a ribosome binding site for translation. The coding portion of the transcripts expressed by the constructs will preferably include a translation initiating codon at the beginning and a termination codon (UAA, UGA or UAG) appropriately positioned at the end of the polypeptide to be translated.

As indicated, the expression vectors will preferably include at least one selectable marker. Such markers include dihydrofolate reductase, G418 or neomycin resistance for eukaryotic cell culture and tetracycline, kanamycin or ampicillin resistance genes for culturing in *E. coli* and other bacteria.



Representative examples of appropriate hosts include, but are not limited to, bacterial cells, such as *E. coli*, *Streptomyces* and *Salmonella typhimurium* cells; fungal cells, such as yeast cells; insect cells such as *Drosophila* S2 and *Spodoptera* Sf9 cells; animal cells such as CHO, COS, 293 and Bowes melanoma cells; and plant cells. Appropriate culture mediums and conditions for the above-described host cells are known in the art.

Among vectors preferred for use in bacteria include pQE70, pQE60 and pQE-9, available from QIAGEN, Inc., *supra*; Phagescript vectors, Bluescript vectors, pNH8A, pNH16a, pNH18A, pNH46A, available from Stratagene; and ptrc99a, pKK223-3, pKK233-3, pDR540, pRIT5 available from Pharmacia. Among preferred eukaryotic vectors are pWLNEO, pSV2CAT, pOG44, pXT1 and pSG available from Stratagene; and pSVK3, pBPV, pMSG and pSVL available from Pharmacia. Other suitable vectors will be readily apparent to the skilled artisan. Introduction of the construct into the host cell can be effected by calcium phosphate transfection, DEAE-dextran mediated transfection, cationic lipid-mediated transfection, electroporation, transduction, infection or other methods. Such methods are described in many standard laboratory manuals, such as Davis *et al.*, *Basic Methods In Molecular Biology* (1986). The polypeptide may be expressed in a modified form, such as a fusion protein, and may include not only secretion signals, but also additional heterologous functional regions. For instance, a region of additional amino acids, particularly charged amino acids, may be added to the N-terminus of the polypeptide to improve stability and persistence in the host cell, during purification, or during subsequent handling and storage. Also, peptide moieties may be added to the polypeptide to facilitate purification. Such regions may be removed prior to final preparation of the polypeptide. The addition of peptide moieties to polypeptides to engender secretion or excretion, to improve stability and to facilitate purification, among others, are familiar and routine techniques in the art. A preferred fusion protein comprises a heterologous region from immunoglobulin that is useful to stabilize and purify proteins. For example, EP-A-O 464 533 (Canadian counterpart 2045869) discloses fusion proteins comprising various portions of constant region of immunoglobulin molecules together with another human protein or part thereof. In many cases, the Fc part in a fusion protein is thoroughly advantageous for use in therapy and diagnosis and thus results, for example, in improved pharmacokinetic properties (EP-A 0232 262). On the

other hand, for some uses it would be desirable to be able to delete the Fc part after the fusion protein has been expressed, detected and purified in the advantageous manner described. This is the case when Fc portion proves to be a hindrance to use in therapy and diagnosis, for example when the fusion protein is to be used as antigen for immunizations. In drug discovery, for example, human proteins, such as hIL-5, have been fused with Fc portions for the purpose of high-throughput screening assays to identify antagonists of hIL-5. See, D. Bennett *et al.*, *J. Molecular Recognition* 8:52-58 (1995) and K. Johanson *et al.*, *J. Biol. Chem.* 270:9459-9471 (1995).

The TNFR proteins can be recovered and purified from recombinant cell cultures by well-known methods including ammonium sulfate or ethanol precipitation, acid extraction, anion or cation exchange chromatography, phosphocellulose chromatography, hydrophobic interaction chromatography, affinity chromatography, hydroxylapatite chromatography and lectin chromatography. Most preferably, high performance liquid chromatography ("HPLC") is employed for purification. Polypeptides of the present invention include: products purified from natural sources, including bodily fluids, tissues and cells, whether directly isolated or cultured; products of chemical synthetic procedures; and products produced by recombinant techniques from a prokaryotic or eukaryotic host, including, for example, bacterial, yeast, higher plant, insect and mammalian cells. Depending upon the host employed in a recombinant production procedure, the polypeptides of the present invention may be glycosylated or may be non-glycosylated. In addition, polypeptides of the invention may also include an initial modified methionine residue, in some cases as a result of host-mediated processes.

### ***Polypeptides and Fragments***

The invention further provides isolated TNFR polypeptides having the amino acid sequences encoded by the deposited cDNAs, or the amino acid

sequences in SEQ ID NOS:2, 4 and 6, or a peptide or polypeptide comprising a portion of the above polypeptides.

#### ***Variant and Mutant Polypeptides***

To improve or alter the characteristics of a TNFR polypeptide, protein engineering may be employed. Recombinant DNA technology known to those skilled in the art can be used to create novel mutant proteins or "muteins" including single or multiple amino acid substitutions, deletions, additions or fusion proteins. Such modified polypeptides can show, e.g., enhanced activity or increased stability. In addition, they may be purified in higher yields and show better solubility than the corresponding natural polypeptide, at least under certain purification and storage conditions.

#### ***N-Terminal and C-Terminal Deletion Mutants***

For instance, for many proteins, including the extracellular domain of a membrane associated protein or the mature form(s) of a secreted protein, it is known in the art that one or more amino acids may be deleted from the N-terminus or C-terminus without substantial loss of biological function. For instance, Ron et al., *J. Biol. Chem.*, 268:2984-2988 (1993) reported modified KGF proteins that had heparin binding activity even if 3, 8, or 27 amino-terminal amino acid residues were missing. In the present case, since the proteins of the invention are members of the TNFR polypeptide family, deletions of N-terminal amino acids up to the Cysteine at position 53 of SEQ ID NO:2 (TNFR-5), and 49 of SEQ ID NOS:4 and 6 (TNFR-6 $\alpha$  and -6 $\beta$ ) may retain some biological activity such as regulation of proliferation and apoptosis of lymphoid cells. Polypeptides having further N-terminal deletions including the C53 residue in SEQ ID NO:2, or the C49 residue in SEQ ID NOS:4 and 6, would not be expected to retain such biological activities because it is known that these residues in a TNFR-related polypeptide are required for forming a disulfide bridge to provide structural stability which is needed for receptor binding and signal transduction.

However, even if deletion of one or more amino acids from the N-terminus of a protein results in modification of loss of one or more biological functions of the protein, other biological activities may still be retained. Thus, the ability of the shortened protein to induce and/or bind to antibodies which recognize the complete or extracellular domain of the TNFR protein generally will be retained when less than the majority of the residues of the complete protein or extracellular domain are removed from the N-terminus. Whether a particular polypeptide lacking N-terminal residues of a complete protein retains such immunologic activities can readily be determined by routine methods described herein and otherwise known in the art.

Accordingly, the present invention further provides polypeptides having one or more residues deleted from the amino terminus of the amino acid sequence of the TNFR shown in SEQ ID NOS:2, 4 and 6, up to the cysteine residue in each which are at position numbers 53, 49 and 49, respectively, and polynucleotides encoding such polypeptides. In particular, the present invention provides TNFk-5 polypeptides comprising the amino acid sequence of residues m-259 of SEQ ID NO:2 where m is an integer in the range of 1-53 where 53 is the position of the first cysteine residue from the N-terminus of the complete TNFR-5 polypeptide (shown in SEQ ID NO:2) believed to be required for activity of the TNFR-5 protein; and TNFR-6 $\alpha$  and -6 $\beta$  polypeptides comprising the amino acid sequence of residues n-300 and 170 of SEQ ID NOS:4 and 6, respectively where n is an integer in the range of 1-49 where 49 is the position of the first cysteine residue from the N-terminus of the complete TNFR-6 $\alpha$  and -6 $\beta$  polypeptides (shown in SEQ ID NOS:4 and 6, respectively) believed to be required for activity of the TNFR-6 $\alpha$  and -6 $\beta$  proteins.

More in particular, the invention provides polynucleotides encoding polypeptides having the amino acid sequence of residues: 1-259, 2-259, 3-259, 4-259, 5-259, 6-259, 7-259, 8-259, 9-259, 10-259, 11-259, 12-259, 13-259, 14-259, 15-259, 16-259, 17-259, 18-259, 19-259, 20-259, 21-259, 22-259, 23-

259, 24-259, 25-259, 26-259, 27-259, 28-259, 29-259, 30-259, 31-259, 32-  
259, 33-259, 34-259, 35-259, 36-259, 37-259, 38-259, 39-259, 40-259, 41-  
259, 42-259, 43-259, 44-259, 45-259, 46-259, 47-259, 48-259, 49-259, 50-  
259, 51-259, 52-259, and 53-259 of SEQ ID NO:2; 1-300, 2-300, 3-300, 4-  
5 300, 5-300, 6-300, 7-300, 8-300, 9-300, 10-300, 11-300, 12-300, 13-300, 14-  
300, 15-300, 16-300, 17-300, 18-300, 19-300, 20-300, 21-300, 22-300, 23-  
300, 24-300, 25-300, 26-300, 27-300, 28-300, 29-300, 30-300, 31-300, 32-  
300, 33-300, 34-300, 35-300, 36-300, 37-300, 38-300, 39-300, 40-300, 41-  
300, 42-300, 43-300, 44-300, 45-300, 46-300, 47-300, 48-300, and 49-300 of  
10 SEQ ID NO:4; and 1-170, 2-170, 3-170, 4-170, 5-170, 6-170, 7-170, 8-170, 9-  
170, 10-170, 11-170, 12-170, 13-170, 14-170, 15-170, 16-170, 17-170, 18-  
170, 19-170, 20-170, 21-170, 22-170, 23-170, 24-170, 25-170, 26-170, 27-  
170, 28-170, 29-170, 30-170, 31-170, 32-170, 33-170, 34-170, 35-170, 36-  
170, 37-170, 38-170, 39-170, 40-170, 41-170, 42-170, 43-170, 44-170, 45-  
15 170, 46-170, 47-170, 48-170, and 49-170 of SEQ ID NO:6. Polynucleotides  
encoding these polypeptides also are provided.

Similarly, many examples of biologically functional C-terminal deletion  
mutants are known. For instance, interferon gamma shows up to ten times  
higher activities by deleting 8-10 amino acid residues from the carboxy  
terminus of the protein (Döbeli et al., *J. Biotechnology* 7:199-216 (1988)). In  
20 the present case, since the protein of the invention is a member of the TNFR  
polypeptide family, deletions of C-terminal amino acids up to the cysteine at  
position 149, 193 and 132 of SEQ ID NOS:2, 4 and 6, respectively, may  
retain some biological activity such as regulation of proliferation and apoptosis  
25 of lymphoid cells. Polypeptides having further C-terminal deletions including  
the cysteines at positions 149, 193, and 132 of SEQ ID NOS:2, 4 and 6,  
respectively, would not be expected to retain such biological activities because  
it is known that these residues in TNF receptor-related polypeptides are  
required for forming disulfide bridges to provide structural stability which is  
30 needed for receptor binding.

However, even if deletion of one or more amino acids from the C-terminus of a protein results in modification or loss of one or more biological functions of the protein, other biological activities may still be retained. Thus, the ability of the shortened protein to induce and/or bind to antibodies which recognize the complete or mature form of the protein generally will be retained when less than the majority of the residues of the complete or mature form protein are removed from the C-terminus. Whether a particular polypeptide lacking C-terminal residues of a complete protein retains such immunologic activities can readily be determined by routine methods described herein and otherwise known in the art.

Accordingly, the present invention further provides polypeptides having one or more residues from the carboxy terminus of the amino acid sequence of TNFR-5, -6 $\alpha$  and -6 $\beta$  shown in SEQ ID NOS:2, 4 and 6 up to the cysteine at position 149, 193 and 132 of SEQ ID NOS:2, 4 and 6, respectively, and polynucleotides encoding such polypeptides. In particular, the present invention provides polypeptides having the amino acid sequence of residues 1-x, 1-y, and 1-z of the amino acid sequence in SEQ ID NOS:2, 4 and 6, respectively, where x is any integer in the range of 149-259, where y is any integer in the range of 193-300, and z is any integer in the range of 132-170. Polynucleotides encoding these polypeptides also are provided.

The invention also provides polypeptides having one or more amino acids deleted from both the amino and the carboxyl termini, which may be described generally as having residues m-x of SEQ ID NO:2, n-y of SEQ ID NO:4, and n-z of SEQ ID NO:6 where m, n, x, y and z are integers as described above.

Also included are a nucleotide sequence encoding a polypeptide consisting of a portion of a complete TNFR amino acid sequence encoded by a cDNA clone contained in ATCC Deposit No. 97798, 97810, or 97809, where this portion excludes from 1 to about 49, 53, or 53 amino acids from the amino terminus of the complete amino acid sequence encoded by the cDNA clone

contained in ATCC Deposit No. 97798, 97810, and 97809, respectively, or from 1 to about 110, 107, or 58 amino acids from the carboxy terminus of the complete amino acid sequence encoded by the cDNA clone contained in ATCC Deposit No. 97798, 97810 and 97809, respectively, or any combination of the above amino terminal and carboxy terminal deletions, of the complete amino acid sequence encoded by the cDNA clone contained in ATCC Deposit No. 97798, 97810 or 97809. Polynucleotides encoding all of the above deletion mutant polypeptide forms also are provided.

#### *Other Mutants*

In addition to terminal deletion forms of the protein discussed above, it also will be recognized by one of ordinary skill in the art that some amino acid sequences of the TNFR polypeptides can be varied without significant effect on the structure or function of the proteins. If such differences in sequence are contemplated, it should be remembered that there will be critical areas on the protein which determine activity.

Thus, the invention further includes variations of the TNFR polypeptides which show substantial TNFR polypeptide activity or which include regions of TNFR protein such as the protein portions discussed below. Such mutants include deletions, insertions, inversions, repeats, and type substitutions selected according to general rules known in the art so as have little effect on activity. For example, guidance concerning how to make phenotypically silent amino acid substitutions is provided in Bowie, J. U. et al., "Deciphering the Message in Protein Sequences: Tolerance to Amino Acid Substitutions," *Science* 247:1306-1310 (1990), wherein the authors indicate that there are two main approaches for studying the tolerance of an amino acid sequence to change. The first method relies on the process of evolution, in which mutations are either accepted or rejected by natural selection. The second approach uses genetic engineering to introduce amino acid changes at



specific positions of a cloned gene and selections or screens to identify sequences that maintain functionality.

As the authors state, these studies have revealed that proteins are surprisingly tolerant of amino acid substitutions. The authors further indicate which amino acid changes are likely to be permissive at a certain position of the protein. For example, most buried amino acid residues require nonpolar side chains, whereas few features of surface side chains are generally conserved. Other such phenotypically silent substitutions are described in Bowie, J. U. *al.*, *supra*, and the references cited therein. Typically seen as conservative substitutions are the replacements, one for another, among the aliphatic amino acids Ala, Val, Leu and Ile; interchange of the hydroxyl residues Ser and Thr, exchange of the acidic residues Asp and Glu, substitution between the amide residues Asn and Gln, exchange of the basic residues Lys and Arg and replacements among the aromatic residues Phe, Tyr. Thus, the fragment, derivative or analog of the polypeptide of SEQ ID NO:2, 4 or 6, or that encoded by a deposited cDNA, may be (i) one in which one or more of the amino acid residues are substituted with a conserved or non-conserved amino acid residue (preferably a conserved amino acid residue) and such substituted amino acid residue may or may not be one encoded by the genetic code, or (ii) one in which one or more of the amino acid residues includes a substituent group, or (iii) one in which the mature or soluble extracellular polypeptide is fused with another compound, such as a compound to increase the half-life of the polypeptide (for example, polyethylene glycol), or (iv) one in which the additional amino acids are fused to the above form of the polypeptide, such as an IgG Fc fusion region peptide or leader or secretory sequence or a sequence which is employed for purification of the above form of the polypeptide or a proprotein sequence. Such fragments, derivatives and analogs are deemed to be within the scope of those skilled in the art from the teachings herein.

Thus, the TNFR of the present invention may include one or more amino acid substitutions, deletions or additions, either from natural mutations

human manipulation. As indicated, changes are preferably of a minor nature, such as conservative amino acid substitutions that do not significantly affect the folding or activity of the protein (see Table 1).

TABLE 1. Conservative Amino Acid Substitutions.

Aromatic	Phenylalanine Tryptophan Tyrosine
Hydrophobic	Leucine Isoleucine Valine
Polar	Glutamine Asparagine
Basic	Arginine Lysine Histidine
Acidic	Aspartic Acid Glutamic Acid
Small	Alanine Serine Threonine Methionine Glycine

Amino acids in the TNFR proteins of the present invention that are essential for function can be identified by methods known in the art, such as site-directed mutagenesis or alanine-scanning mutagenesis (Cunningham and Wells, *Science* 244:1081-1085 (1989)). The latter procedure introduces single alanine mutations at every residue in the molecule. The resulting mutant molecules are then tested for biological activity such as receptor binding or *in vitro* or *in vitro* proliferative activity.

Of special interest are substitutions of charged amino acids with other charged or neutral amino acids which may produce proteins with highly desirable improved characteristics, such as less aggregation. Aggregation may not only reduce activity but also be problematic when preparing pharmaceutical formulations, because aggregates can be immunogenic (Pinckard *et al.*, *Clin. Exp. Immunol.* 2:331-340 (1967); Robbins *et al.*, *Diabetes* 36: 838-

845 (1987); Cleland *et al.*, *Crit. Rev. Therapeutic Drug Carrier Systems* 10:307-377 (1993).

Replacement of amino acids can also change the selectivity of the binding of a ligand to cell surface receptors. For example, Ostade *et al.*, *Nature* 361:266-268 (1993) describes certain mutations resulting in selective binding of TNF- $\alpha$  to only one of the two known types of TNF receptors. Sites that are critical for ligand-receptor binding can also be determined by structural analysis such as crystallization, nuclear magnetic resonance or photoaffinity labeling (Smith *et al.*, *J. Mol. Biol.* 224:899-904 (1992) and de Vos *et al.* *Science* 255:306-312 (1992)).

Since TNFR-5, -6 $\alpha$  and -6 $\beta$  are members of the TNF receptor-related protein family, to modulate rather than completely eliminate biological activities of TNFR preferably mutations are made in sequences encoding amino acids in the TNFR conserved extracellular domain, more preferably in residues within this region which are not conserved among members of the TNF receptor family. Also forming part of the present invention are isolated polynucleotides comprising nucleic acid sequences which encode the above TNFR mutants.

The polypeptides of the present invention are preferably provided in an isolated form, and preferably are substantially purified. A recombinantly produced version of the TNFR polypeptides can be substantially purified by the one-step method described in Smith and Johnson, *Gene* 67:31-40 (1988). Polypeptides of the invention also can be purified from natural or recombinant sources using anti-TNFR-5, -6 $\alpha$  and -6 $\beta$  antibodies of the invention in methods which are well known in the art of protein purification.

The invention further provides isolated TNFR polypeptides comprising an amino acid sequence selected from the group consisting of: (a) the amino acid sequence of a full-length TNFR polypeptide having the complete amino acid sequence shown in SEQ ID NO:2, 4 or 6 or as encoded by a cDNA clone contained in ATCC Deposit No. 97798, 97810 or 97809; (b) the amino acid sequence of a mature TNFR polypeptide having the amino acid sequence at positions 27-259 in SEQ ID NO:2, 31-300 in SEQ ID NO:4, or 31-170 in SEQ ID NO:6, or as encoded by a cDNA clone contained in ATCC Deposit No. 97798, 97810 or 97809; or (c) the amino acid sequence of a soluble extracellular domain of a TNFR polypeptide having the amino acid sequence at positions 27 to 240 in SEQ ID NO:2, 31 to 283 in SEQ ID NO:4, or 31 to 166 in SEQ ID NO:6, or as encoded by the cDNA clone contained in ATCC Deposit No. 97798, 97810 or 97809.

Further polypeptides of the present invention include polypeptides which have at least 90% similarity, more preferably at least 95% similarity, and still more preferably at least 96%, 97%, 98% or 99% similarity to those described above. The polypeptides of the invention also comprise those which are at least 80% identical, more preferably at least 90% or 95% identical, still more preferably at least 96%, 97%, 98% or 99% identical to the polypeptide encoded by the deposited cDNA or to the polypeptide of SEQ ID NO:2, 4 or 6, and also include portions of such polypeptides with at least 30 amino acids and more preferably at least 50 amino acids.

By "% similarity" for two polypeptides is intended a similarity score produced by comparing the amino acid sequences of the two polypeptides using the Bestfit program (Wisconsin Sequence Analysis Package, Version 8 for Unix, Genetics Computer Group, University Research Park, 575 Science Drive, Madison, WI 53711) and the default settings for determining similarity. Bestfit uses the local homology algorithm of Smith and Waterman (Advances in Applied Mathematics 2:482-489, 1981) to find the best segment of similarity between two sequences.

By a polypeptide having an amino acid sequence at least, for example, 95% "identical" to a reference amino acid sequence of a TNFR polypeptide is intended that the amino acid sequence of the polypeptide is identical to the reference sequence except that the polypeptide sequence may include up to five amino acid alterations per each 100 amino acids of the reference amino acid of the TNFR polypeptide. In other words, to obtain a polypeptide having an amino acid sequence at least 95% identical to a reference amino acid sequence, up to 5% of the amino acid residues in the reference sequence may be deleted or substituted with another amino acid, or a number of amino acids up to 5% of the total amino acid residues in the reference sequence may be inserted into the reference sequence. These alterations of the reference sequence may occur at the amino or carboxy terminal positions of the reference amino acid sequence or anywhere between those terminal positions, interspersed either individually among residues in the reference sequence or in one or more contiguous groups within the reference sequence.

As a practical matter, whether any particular polypeptide is at least 90%, 95%, 96%, 97%, 98% or 99% identical to, for instance, the amino acid sequence shown in SEQ ID NO:2, 4 or 6, or to an amino acid sequence encoded by a deposited cDNA clone can be determined conventionally using known computer programs such the Bestfit program (Wisconsin Sequence Analysis Package, Version 8 for Unix, Genetics Computer Group, University Research Park, 575 Science Drive, Madison, WI 53711). When using Bestfit or any other sequence alignment program to determine whether a particular sequence is, for instance, 95% identical to a reference sequence according to the present invention, the parameters are set, of course, such that the percentage of identity is calculated over the full length of the reference amino acid sequence and that gaps in homology of up to 5% of the total number of amino acid residues in the reference sequence are allowed.

The polypeptide of the present invention could be used as a molecular weight marker on SDS-PAGE gels or on molecular sieve gel filtration columns using methods well known to those of skill in the art.

As described in detail below, the polypeptides of the present invention can also be used to raise polyclonal and monoclonal antibodies, which are useful in assays for detecting TNFR protein expression as described below or as agonists and antagonists capable of enhancing or inhibiting TNFR protein function. Further, such polypeptides can be used in the yeast two-hybrid system to "capture" TNFR protein binding proteins which are also candidate agonists and antagonists according to the present invention. The yeast two hybrid system is described in Fields and Song, *Nature* 340:245-246 (1989).

#### *Epitope-Bearing Portions*

In another aspect, the invention provides a peptide or polypeptide comprising an epitope-bearing portion of a polypeptide of the invention. The epitope of this polypeptide portion is an immunogenic or antigenic epitope of a polypeptide of the invention. An "immunogenic epitope" is defined as a part of a protein that elicits an antibody response when the whole protein is the immunogen. On the other hand, a region of a protein molecule to which an antibody can bind is defined as an "antigenic epitope." The number of immunogenic epitopes of a protein generally is less than the number of antigenic epitopes. See, for instance, Geysen *et al.*, *Proc. Natl. Acad. Sci. USA* 81:3998-4002 (1983).

As to the selection of peptides or polypeptides bearing an antigenic epitope (i.e., that contain a region of a protein molecule to which an antibody can bind), it is well known in that art that relatively short synthetic peptides that mimic part of a protein sequence are routinely capable of eliciting an antiserum that reacts with the partially mimicked protein. See, for instance, Sutcliffe, J. G., Shinnick, T. M., Green, N. and Learner, R. A. (1983) "Antibodies that react with predetermined sites on proteins," *Science*, 219:660-666. Peptides capable of eliciting protein-reactive sera are frequently

represented in the primary sequence of a protein, can be characterized by a set of simple chemical rules, and are confined neither to immunodominant regions of intact proteins (i.e., immunogenic epitopes) nor to the amino or carboxyl terminals. Antigenic epitope-bearing peptides and polypeptides of the invention are therefore useful to raise antibodies, including monoclonal antibodies, that bind specifically to a polypeptide of the invention. See, for instance, Wilson *et al.*, *Cell* 37:767-778 (1984) at 777.

Antigenic epitope-bearing peptides and polypeptides of the invention preferably contain a sequence of at least seven, more preferably at least nine and most preferably between about 15 to about 30 amino acids contained within the amino acid sequence of a polypeptide of the invention.

Non-limiting examples of antigenic polypeptides or peptides that can be used to generate TNFR-specific antibodies include: a polypeptide comprising amino acid residues from about Gln-42 to about Glu-52, from about His-58 to about Cys-66, from about Pro-68 to about Thr-76, from about Ser-79 to about Cys-85, from about Cys-91 to about Thr-102, from about Gln-110 to about Pro-122, from about Arg-126 to about Val-136, and from about Thr-142 to about Glu-148 in SEQ ID NO:2; from about Ala-31 to about Thr-46, from about Phe-57 to about Thr-117, from about Cys-132 to about Thr-175, from about Gly-185 to about Thr-194, from about Val-205 to about Asp-217, from about Pro-239 to about Leu-264, and from about Ala-283 to about Pro-298 in SEQ ID NO:4; and from about Ala-31 to about Thr-46, from about Phe-57 to about Gln-80, from about Glu-86 to about His-106, from about Thr-108 to about Phe-119, from about His-129 to about Val-138, and from about Gly-142 to about Pro-166 in SEQ ID NO:6. These polypeptide fragments have been determined to bear antigenic epitopes of the TNFR-5, -6 $\alpha$  and -6 $\beta$  polypeptides respectively, by the analysis of the Jameson-Wolf antigenic index, as shown in Figures 5, 6 and 7, above.

The epitope-bearing peptides and polypeptides of the invention may be produced by any conventional means. See, e.g., Houghten, R. A. (1985)



"General method for the rapid solid-phase synthesis of large numbers of peptides: specificity of antigen-antibody interaction at the level of individual amino acids." *Proc. Natl. Acad. Sci. USA* 82:5131-5135; this "Simultaneous Multiple Peptide Synthesis (SMPS)" process is further described in U.S. Patent No. 4,631,211 to Houghten *et al.* (1986).

Epitope-bearing peptides and polypeptides of the invention are used to induce antibodies according to methods well known in the art. See, for instance, Sutcliffe *et al.*, *supra*; Wilson *et al.*, *supra*; Chow, M. *et al.*, *Proc. Natl. Acad. Sci. USA* 82:910-914; and Bittle, F. J. *et al.*, *J. Gen. Virol.* 66:2347-2354 (1985). Immunogenic epitope-bearing peptides of the invention, i.e., those parts of a protein that elicit an antibody response when the whole protein is the immunogen, are identified according to methods known in the art. See, for instance, Geysen *et al.*, *supra*. Further still, U.S. Patent No. 5,194,392 to Geysen (1990) describes a general method of detecting or determining the sequence of monomers (amino acids or other compounds) which is a topological equivalent of the epitope (i.e., a "mimotope") which is complementary to a particular paratope (antigen binding site) of an antibody of interest. More generally, U.S. Patent No. 4,433,092 to Geysen (1989) describes a method of detecting or determining a sequence of monomers which is a topographical equivalent of a ligand which is complementary to the ligand binding site of a particular receptor of interest. Similarly, U.S. Patent No. 5,480,971 to Houghten, R. A. *et al.* (1996) on Feralkylated Oligopeptide Mixtures discloses linear C1-C7-alkyl peralkylated oligopeptides and sets and libraries of such peptides, as well as methods for using such oligopeptide sets and libraries for determining the sequence of a peralkylated oligopeptide that preferentially binds to an acceptor molecule of interest. Thus, non-peptide analogs of the epitope-bearing peptides of the invention also can be made routinely by these methods.

### *Fusion Proteins*

As one of skill in the art will appreciate, TNFR polypeptides of the present invention and the epitope-bearing fragments thereof described above can be combined with parts of the constant domain of immunoglobulins (IgG), resulting in chimeric polypeptides. These fusion proteins facilitate purification and show an increased half-life in vivo. This has been shown, e.g., for chimeric proteins consisting of the first two domains of the human CD4-polypeptide and various domains of the constant regions of the heavy or light chains of mammalian immunoglobulins (EP A 394,827; Traunecker et al., Nature 331:84-86 (1988)). Fusion proteins that have a disulfide-linked dimeric structure due to the IgG part can also be more efficient in binding and neutralizing other molecules than the monomeric TNFR protein or protein fragment alone (Fountoulakis et al., *J. Biochem.* 270:3958-3964 (1995)).

### *Antibodies*

TNFR-protein specific antibodies for use in the present invention can be raised against the intact TNFR-5, -6 $\alpha$  and -6 $\beta$  proteins or an antigenic polypeptide fragment thereof, which may be presented together with a carrier protein, such as an albumin, to an animal system (such as rabbit or mouse) or, if it is long enough (at least about 25 amino acids), without a carrier.

As used herein, the term "antibody" (Ab) or "monoclonal antibody" (Mab) is meant to include intact molecules as well as antibody fragments (such as, for example, Fab and F(ab')<sub>2</sub> fragments) which are capable of specifically binding to a TNFR protein. Fab and F(ab')<sub>2</sub> fragments lack the Fc fragment of intact antibody, clear more rapidly from the circulation, and may have less non-specific tissue binding of an intact antibody (Wahl et al., *J. Nucl. Med.* 24:316-325 (1983)). Thus, these fragments are preferred.

The antibodies of the present invention may be prepared by any of a variety of methods. For example, cells expressing the TNFR protein or an antigenic fragment thereof can be administered to an animal in order to induce

the production of sera containing polyclonal antibodies. In a preferred method, a preparation of TNFR protein is prepared and purified to render it substantially free of natural contaminants. Such a preparation is then introduced into an animal in order to produce polyclonal antisera of greater specific activity.

In the most preferred method, the antibodies of the present invention are monoclonal antibodies. Such monoclonal antibodies can be prepared using hybridoma technology (Köhler *et al.*, *Nature* 256:495 (1975); Köhler *et al.*, *Eur. J. Immunol.* 6:511 (1976); Köhler *et al.*, *Eur. J. Immunol.* 6:292 (1976); Hammerling *et al.*, in: *Monoclonal Antibodies and T-Cell Hybridomas*, Elsevier, N.Y., (1981) pp. 563-681). In general, such procedures involve immunizing an animal (preferably a mouse) with a TNFR protein antigen or, more preferably, with a TNFR protein-expressing cell. Suitable cells can be recognized by their capacity to bind anti-TNFR-5, -6 $\alpha$  or -6 $\beta$  protein antibody. Such cells may be cultured in any suitable tissue culture medium; however, it is preferable to culture cells in Earle's modified Eagle's medium supplemented with 10% fetal bovine serum (inactivated at about 56° C), and supplemented with about 10 g/l of nonessential amino acids, about 1,000 U/ml of penicillin, and about 100  $\mu$ g/ml of streptomycin. The splenocytes of such mice are extracted and fused with a suitable myeloma cell line. Any suitable myeloma cell line may be employed in accordance with the present invention; however, it is preferable to employ the parent myeloma cell line (SP2O), available from the American Type Culture Collection, Rockville, Maryland. After fusion, the resulting hybridoma cells are selectively maintained in HAT medium, and then cloned by limiting dilution as described by Wands *et al.* (Gastroenterology 80:225-232 (1981)). The hybridoma cells obtained through such a selection are then assayed to identify clones which secrete antibodies capable of binding the desired TNFR antigen.

Alternatively, additional antibodies capable of binding to the TNFR antigen may be produced in a two-step procedure through the use of

anti-idiotypic antibodies. Such a method makes use of the fact that antibodies are themselves antigens, and that, therefore, it is possible to obtain an antibody which binds to a second antibody. In accordance with this method, TNFR-protein specific antibodies are used to immunize an animal, preferably a mouse. The splenocytes of such an animal are then used to produce hybridoma cells, and the hybridoma cells are screened to identify clones which produce an antibody whose ability to bind to the TNFR protein-specific antibody can be blocked by the TNFR protein antigen. Such antibodies comprise anti-idiotypic antibodies to the TNFR protein-specific antibody and can be used to immunize an animal to induce formation of further TNFR protein-specific antibodies.

It will be appreciated that Fab and F(ab')<sub>2</sub> and other fragments of the antibodies of the present invention may be used according to the methods disclosed herein. Such fragments are typically produced by proteolytic cleavage, using enzymes such as papain (to produce Fab fragments) or pepsin (to produce F(ab')<sub>2</sub> fragments). Alternatively, TNFR protein-binding fragments can be produced through the application of recombinant DNA technology or through synthetic chemistry.

For *in vivo* use of anti-TNFR in humans, it may be preferable to use "humanized" chimeric monoclonal antibodies. Such antibodies can be produced using genetic constructs derived from hybridoma cells producing the monoclonal antibodies described above. Methods for producing chimeric antibodies are known in the art. See, for review, Morrison, *Science* 229:1202 (1985); Oi et al., *BioTechniques* 4:214 (1986); Cabilly et al., U.S. Patent No. 4,816,567; Taniguchi et al., EP 171496; Morrison et al., EP 173494; Neuberger et al., WO 8601533; Robinson et al., WO 8702671; Boulianne et al., *Nature* 312:643 (1984); Neuberger et al., *Nature* 314:268 (1985).

### *Immune System-Related Disorders*

#### *Diagnosis*

The present inventors have discovered that TNFR-5, -6 $\alpha$  and -6 $\beta$  are expressed in hematopoietic tissues. For a number of immune system-related disorders, substantially altered (increased or decreased) levels of TNFR gene expression can be detected in immune system tissue or other cells or bodily fluids (e.g., sera and plasma) taken from an individual having such a disorder, relative to a "standard" TNFR gene expression level, that is, the TNFR expression level in immune system tissues or bodily fluids from an individual not having the immune system disorder. Thus, the invention provides a diagnostic method useful during diagnosis of an immune system disorder, which involves measuring the expression level of the gene encoding the TNFR protein in immune system tissue or other cells or body fluid from an individual and comparing the measured gene expression level with a standard TNFR gene expression level, whereby an increase or decrease in the gene expression level compared to the standard is indicative of an immune system disorder.

In particular, it is believed that certain tissues in mammals with cancer express significantly reduced levels of the TNFR protein and mRNA encoding the TNFR when compared to a corresponding "standard" level. Further, it is believed that reduced levels of the TNFR protein can be detected in certain body fluids (e.g., sera and plasma) from mammals with such a cancer when compared to sera from mammals of the same species not having the cancer.

Thus, the invention provides a diagnostic method useful during diagnosis of an immune system disorder, including cancers which involves measuring the expression level of the gene encoding the TNFR protein in immune system tissue or other cells or body fluid from an individual and comparing the measured gene expression level with a standard TNFR gene expression level, whereby an increase or decrease in the gene expression level compared to the standard is indicative of an immune system disorder.

Where a diagnosis of a disorder in the immune system including diagnosis of a tumor has already been made according to conventional methods, the present invention is useful as a prognostic indicator, whereby patients exhibiting depressed gene expression will experience a worse clinical outcome relative to patients expressing the gene at a level nearer the standard level.

By "assaying the expression level of the gene encoding a TNFR protein" is intended qualitatively or quantitatively measuring or estimating the level of the TNFR-5, -6 $\alpha$  and/or -6 $\beta$  protein or the level of the mRNA encoding the TNFR-5, -6 $\alpha$  and/or -6 $\beta$  protein in a first biological sample either directly (e.g., by determining or estimating absolute protein level or mRNA level) or relatively (e.g., by comparing to the TNFR protein level or mRNA level in a second biological sample). Preferably, the TNFR protein level or mRNA level in the first biological sample is measured or estimated and compared to a standard TNFR protein level or mRNA level, the standard being taken from a second biological sample obtained from an individual not having the disorder or being determined by averaging levels from a population of individuals not having a disorder of the immune system. As will be appreciated in the art, once standard TNFR protein levels or mRNA levels are known, they can be used repeatedly as a standard for comparison.

By "biological sample" is intended any biological sample obtained from an individual, body fluid, cell line, tissue culture, or other source which contains TNFR protein or mRNA. As indicated, biological samples include body fluids (such as sera, plasma, urine, synovial fluid and spinal fluid) which contain free extracellular domain(s) (or soluble form(s)) of a TNFR protein, immune system tissue, and other tissue sources found to express complete or extracellular domain of a TNFR. Methods for obtaining tissue biopsies and body fluids from mammals are well known in the art. Where the biological sample is to include mRNA, a tissue biopsy is the preferred source.

The invention also contemplates the use of a gene of the present invention for diagnosing mutations in a TNFR gene. For example, if a

mutation is present in one of the genes of the present invention, conditions would result from a lack of production of the receptor polypeptides of the present invention. Further, mutations which enhance receptor polypeptide activity would lead to diseases associated with an over expression of the receptor polypeptide, e.g., endotoxic shock. Mutations in the genes can be detected by comparing the sequence of the defective gene with that of a normal one. Subsequently one can verify that a mutant gene is associated with a disease condition or the susceptibility to a disease condition. That is, a mutant gene which leads to the underexpression of the receptor polypeptides of the present invention would be associated with an inability of TNF to inhibit tumor growth.

Other immune system disorders which may be diagnosed by the foregoing assays include hypersensitivity, allergy, infectious disease, graft-host disease, immunodeficiency, autoimmune diseases and the like.

Individuals carrying mutations in the genes of the present invention may be detected at the DNA level by a variety of techniques. Nucleic acids used for diagnosis may be obtained from a patient's cells, such as from blood, urine, saliva and tissue biopsy among other tissues. The genomic DNA may be used directly for detection or may be amplified enzymatically by using PCR (Saiki et al., Nature, 324:163-166 (1986)) prior to analysis. RNA or cDNA may also be used for the same purpose. As an example, PCR primers complementary to the nucleic acid of the instant invention can be used to identify and analyze mutations in the human genes of the present invention. For example, deletions and insertions can be detected by a change in the size of the amplified product in comparison to the normal genotype. Point mutations can be identified by hybridizing amplified DNA to radiolabeled RNA or alternatively, radiolabeled antisense DNA sequences of the present invention. Perfectly matched sequences can be distinguished from mismatched duplexes by RNase A digestion or by differences in melting temperatures. Such a diagnostic would be particularly useful for prenatal or even neonatal testing.

Sequence differences between the reference gene and "mutants" may be revealed by the direct DNA sequencing method. In addition, cloned DNA segments may be used as probes to detect specific DNA segments. The sensitivity of this method is greatly enhanced when combined with PCR. For example, a sequencing primary used with double stranded PCR product or a single stranded template molecule generated by a modified PCR product. The sequence determination is performed by conventional procedures with radiolabeled nucleotides or by automatic sequencing procedures with fluorescent tags.

Sequence changes at the specific locations may be revealed by nuclease protection assays, such as RNase and S1 protection or the chemical cleavage method (for example, Cotton et al., PNAS, 85:4397-4401 (1985)).

Assaying TNFR protein levels in a biological sample can occur using antibody-based techniques. For example, TNFR protein expression in tissues can be studied with classical immunohistological methods (Jalkanen, M., et al., *J. Cell. Biol.* 101:976-985 (1985); Jalkanen, M., et al., *J. Cell. Biol.* 105:3087-3096 (1987)). Other antibody-based methods useful for detecting TNFR gene expression include immunoassays, such as the enzyme linked immunosorbent assay (ELISA) and the radioimmunoassay (RIA). Suitable antibody assay labels are known in the art and include enzyme labels, such as, glucose oxidase, and radioisotopes, such as iodine ( $^{125}\text{I}$ ,  $^{121}\text{I}$ ), carbon ( $^{14}\text{C}$ ), sulfur ( $^{35}\text{S}$ ), tritium ( $^3\text{H}$ ), indium ( $^{112}\text{In}$ ), and technetium ( $^{99\text{m}}\text{Tc}$ ), and fluorescent labels, such as fluorescein and rhodamine, and biotin.

In addition to assaying TNFR protein levels in a biological sample obtained from an individual, TNFR proteins can also be detected *in vivo* by imaging. Antibody labels or markers for *in vivo* imaging of TNFR proteins include those detectable by X-radiography, NMR or ESR. For X-radiography, suitable labels include radioisotopes such as barium or cesium, which emit detectable radiation but are not overtly harmful to the subject. Suitable markers for NMR and ESR include those with a detectable characteristic spin.



such as deuterium, which may be incorporated into the antibody by labeling of nutrients for the relevant hybridoma.

A TNFR-specific antibody or antibody fragment which has been labeled with an appropriate detectable imaging moiety, such as a radioisotope (for example,  $^{131}\text{I}$ ,  $^{112}\text{In}$ ,  $^{99\text{m}}\text{Tc}$ ), a radio-opaque substance, or a material detectable by nuclear magnetic resonance, is introduced (for example, parenterally, subcutaneously or intraperitoneally) into the mammal to be examined for immune system disorder. It will be understood in the art that the size of the subject and the imaging system used will determine the quantity of imaging moiety needed to produce diagnostic images. In the case of a radioisotope moiety, for a human subject, the quantity of radioactivity injected will normally range from about 5 to 20 millicuries of  $^{99\text{m}}\text{Tc}$ . The labeled antibody or antibody fragment will then preferentially accumulate at the location of cells which contain TNFR protein. *In vivo* tumor imaging is described in S.W. Burchiel et al., "Immunopharmacokinetics of Radiolabeled Antibodies and Their Fragments" (Chapter 13 in *Tumor Imaging: The Radiochemical Detection of Cancer*, S.W. Burchiel and B. A. Rhodes, eds., Masson Publishing Inc. (1982)).

#### *Treatment*

The Tumor Necrosis Factor (TNF) family ligands are known to be among the most pleiotropic cytokines, inducing a large number of cellular responses, including cytotoxicity, anti-viral activity, immunoregulatory activities, and the transcriptional regulation of several genes (Goeddel, D.V. et al., "Tumor Necrosis Factors: Gene Structure and Biological Activities," *Symp. Quant. Biol.* 51:597-609 (1986). Cold Spring Harbor; Beutler, B., and Cerami, A., *Annu. Rev. Biochem.* 57:505-518 (1988); Old, L.J., *Sci. Am.* 258:59-75 (1988); Fiers, W., *FEBS Lett.* 285:199-224 (1991)). The TNF-family ligands induce such various cellular responses by binding to TNF-family receptors. Cells which express a TNFR polypeptide and have a potent cellular response to TNFR-5, -6 $\alpha$  and -6 $\beta$  ligands include lymphocytes, endothelial cells, keratinocytes, and prostate tissue. By "a cellular response to a TNF-family ligand" is intended any genotypic, phenotypic, and/or

morphologic change to a cell, cell line, tissue, tissue culture or patient that is induced by a TNF-family ligand. As indicated, such cellular responses include not only normal physiological responses to TNF-family ligands, but also diseases associated with increased apoptosis or the inhibition of apoptosis.

5 Diseases associated with increased cell survival, or the inhibition of apoptosis, include cancers (such as follicular lymphomas, carcinomas with p53 mutations, and hormone-dependent tumors, such as breast cancer, prostate cancer, Kaposi's sarcoma and ovarian cancer); autoimmune disorders (such as systemic lupus erythematosus and immune-related glomerulonephritis  
10 rheumatoid arthritis) and viral infections (such as herpes viruses, pox viruses and adenoviruses), information graft v. host disease, acute graft rejection, and chronic graft rejection. Diseases associated with increased apoptosis include AIDS; neurodegenerative disorders (such as Alzheimer's disease, Parkinson's  
15 disease, Amyotrophic lateral sclerosis, Retinitis pigmentosa, Cerebellar degeneration); myelodysplastic syndromes (such as aplastic anemia), ischemic injury (such as that caused by myocardial infarction, stroke and reperfusion injury), toxin-induced liver disease (such as that caused by alcohol), septic shock, cachexia and anorexia.

20 Thus, in one aspect, the present invention is directed to a method for enhancing apoptosis induced by a TNF-family ligand, which involves administering to a cell which expresses the TNFR polypeptide an effective amount of TNFR polypeptide, analog or an agonist capable of increasing TNFR mediated signaling. Preferably, TNFR mediated signaling is increased to treat a disease wherein decreased apoptosis is exhibited. Antagonist can  
25 include soluble forms of TNFR and monoclonal antibodies directed against the TNFR polypeptide.

30 By "agonist" is intended naturally occurring and synthetic compounds capable of enhancing or potentiating apoptosis. By "antagonist" is intended naturally occurring and synthetic compounds capable of inhibiting apoptosis. Whether any candidate "agonist" or "antagonist" of the present invention can enhance or inhibit apoptosis can be determined using art-known TNF-family ligand/receptor cellular response assays, including those described in more detail below.

35 One such screening procedure involves the use of melanophores which are transfected to express the receptor of the present invention. Such a screening technique is described in PCT WO 92 01810, published February 6, 1992. Such an assay may be employed, for example, for screening for a compound which inhibits (or enhances) activation of the receptor polypeptide

of the present invention by contacting the melanophore cells which encode the receptor with both a TNF-family ligand and the candidate antagonist (or agonist). Inhibition or enhancement of the signal generated by the ligand indicates that the compound is an antagonist or agonist of the ligand/receptor signaling pathway.

Other screening techniques include the use of cells which express the receptor (for example, transfected CHO cells) in a system which measures extracellular pH changes caused by receptor activation, for example, as described in *Science* 246:181-296 (October 1989). For example, compounds may be contacted with a cell which expresses the receptor polypeptide of the present invention and a second messenger response, e.g., signal transduction or pH changes, may be measured to determine whether the potential compound activates or inhibits the receptor.

Another such screening technique involves introducing RNA encoding the receptor into *Xenopus* oocytes to transiently express the receptor. The receptor oocytes may then be contacted with the receptor ligand and a compound to be screened, followed by detection of inhibition or activation of a calcium signal in the case of screening for compounds which are thought to inhibit activation of the receptor.

Another screening technique involves expressing in cells a construct wherein the receptor is linked to a phospholipase C or D. Such cells include endothelial cells, smooth muscle cells, embryonic kidney cells, etc. The screening may be accomplished as hereinabove described by detecting activation of the receptor or inhibition of activation of the receptor from the phospholipase signal.

Another method involves screening for compounds which inhibit activation of the receptor polypeptide of the present invention antagonists by determining inhibition of binding of labeled ligand to cells which have the receptor on the surface thereof. Such a method involves transfecting a eukaryotic cell with DNA encoding the receptor such that the cell expresses the receptor on its surface and contacting the cell with a compound in the presence of a labeled form of a known ligand. The ligand can be labeled, e.g., by radioactivity. The amount of labeled ligand bound to the receptors is measured, e.g., by measuring radioactivity of the receptors. If the compound binds to the receptor as determined by a reduction of labeled ligand which binds to the receptors, the binding of labeled ligand to the receptor is inhibited.

Further screening assays for agonist and antagonist of the present invention are described in Tartaglia, L.A., and Goeddel, D.V., *J. Biol. Chem.* 267(7):4304-4307(1992).

Thus, in a further aspect, a screening method is provided for determining whether a candidate agonist or antagonist is capable of enhancing or inhibiting a cellular response to a TNF-family ligand. The method involves contacting cells which express the TNFR polypeptide with a candidate compound and a TNF-family ligand, assaying a cellular response, and comparing the cellular response to a standard cellular response, the standard being assayed when contact is made with the ligand in absence of the candidate compound, whereby an increased cellular response over the standard indicates that the candidate compound is an agonist of the ligand/receptor signaling pathway and a decreased cellular response compared to the standard indicates that the candidate compound is an antagonist of the ligand/receptor signaling pathway. By "assaying a cellular response" is intended qualitatively or quantitatively measuring a cellular response to a candidate compound and/or a TNF-family ligand (e.g., determining or estimating an increase or decrease in T cell proliferation or tritiated thymidine labeling). By the invention, a cell expressing the TNFR polypeptide can be contacted with either an endogenous or exogenously administered TNF-family ligand.

Agonist according to the present invention include naturally occurring and synthetic compounds such as, for example, TNF family ligand peptide fragments, transforming growth factor, neurotransmitters (such as glutamate, dopamine, *N*-methyl-D-aspartate), tumor suppressors (p53), cytolytic T cells and antimetabolites. Preferred agonist include chemotherapeutic drugs such as, for example, cisplatin, doxorubicin, bleomycin, cytosine arabinoside, nitrogen mustard, methotrexate and vincristine. Others include ethanol and  $\beta$ -amyloid peptide. (*Science* 267:1457-1458 (1995)). Further preferred agonist include polyclonal and monoclonal antibodies raised against the TNFR polypeptide, or a fragment thereof. Such agonist antibodies raised against a TNF-family receptor are disclosed in Tartaglia, L.A., et al., *Proc. Natl. Acad. Sci. USA* 88:9292-9296 (1991); and Tartaglia, L.A., and Goeddel, D.V., *J. Biol. Chem.* 267 (7):4304-4307 (1992) See, also, PCT Application WO 94/09137.

Antagonist according to the present invention include naturally occurring and synthetic compounds such as, for example, the CD40 ligand, neutral amino acids, zinc, estrogen, androgens, viral genes (such as Adenovirus *E1B*, Baculovirus *p35* and *IAP*, Cowpox virus *crmA*, Epstein-Barr virus *BIIRF1*, *LMP-1*, African swine fever virus *LMW5-HL*, and Herpesvirus *yl*

34.5), calpain inhibitors, cysteine protease inhibitors, and tumor promoters (such as PMA, Phenobarbital, and  $\gamma$ -Hexachlorocyclohexane). Other antagonists include polyclonal and monoclonal antagonist antibodies raised against the TNFR polypeptides or a fragment thereof. Such antagonist antibodies raised against a TNF-family receptor are described in Tartaglia, L.A., and Goeddel, D.V., *J. Biol. Chem.* 267(7):4304-4307 (1992) and Tartaglia, L.A. *et al.*, *Cell* 73:213-216 (1993). See, also, PCT Application WO 94/09137.

Other potential antagonists include antisense molecules. Antisense technology can be used to control gene expression through antisense DNA or RNA or through triple-helix formation. Antisense techniques are discussed, for example, in Okano, *Neurochem.* 56:560 (1991); *Oligodeoxynucleotides as Antisense Inhibitors of Gene Expression*, CRC Press, Boca Raton, FL (1988). Triple helix formation is discussed in, for instance Lee *et al.*, *Nucleic Acids Research* 6:3073 (1979); Cooney *et al.*, *Science* 241:456 (1988); and Dervan *et al.*, *Science* 251:1360 (1991). The methods are based on binding of a polynucleotide to a complementary DNA or RNA.

For example, the 5' coding portion of a polynucleotide that encodes the mature polypeptide of the present invention may be used to design an antisense RNA oligonucleotide of from about 10 to 40 base pairs in length. A DNA oligonucleotide is designed to be complementary to a region of the gene involved in transcription thereby preventing transcription and the production of the receptor. The antisense RNA oligonucleotide hybridizes to the mRNA *in vivo* and blocks translation of the mRNA molecule into receptor polypeptide. The oligonucleotides described above can also be delivered to cells such that the antisense RNA or DNA may be expressed *in vivo* to inhibit production of the receptor.

Further antagonist according to the present invention include soluble forms of TNFR, i.e., TNFR fragments that include the ligand binding domain from the extracellular region of the full length receptor. Such soluble forms of the receptor, which may be naturally occurring or synthetic, antagonize TNFR mediated signaling by competing with the cell surface TNFR for binding to TNF-family ligands. Thus, soluble forms of the receptor that include the ligand binding domain are novel cytokines capable of inhibiting tumor necrosis induced by TNF-family ligands. Other such cytokines are known in the art and include Fas B (a soluble form of the mouse Fas receptor) that acts physiologically to limit apoptosis induced by Fas ligand (Hughes, D.P. and Crispe, I.N., *J. Exp. Med.* 182:1395-1401 (1995)).

As indicated polyclonal and monoclonal antibody agonist or antagonist according to the present invention can be raised according to the methods disclosed in Tartaglia, L.A., and Goeddel, D.V., *J. Biol. Chem.* 267(7):4304-4307(1992); Tartaglia, L.A. *et al.*, *Cell* 73:213-216 (1993), and PCT Application WO 94/09137. The term "antibody" (Ab) or "monoclonal antibody" (mAb) as used herein is meant to include intact molecules as well as fragments thereof (such as, for example, Fab and F(ab')<sub>2</sub> fragments) which are capable of binding an antigen. Fab and F(ab')<sub>2</sub> fragments lack the Fc fragment of intact antibody, clear more rapidly from the circulation, and may have less non-specific tissue binding of an intact antibody (Wahl *et al.*, *J. Nucl. Med.* 24:316-325 (1983)).

Antibodies according to the present invention may be prepared by any of a variety of methods described above, and known in the art

Proteins and other compounds which bind the extracellular domains are also candidate agonist and antagonist according to the present invention. Such binding compounds can be "captured" using the yeast two-hybrid system (Fields and Song, *Nature* 340:245-246 (1989)). A modified version of the yeast two-hybrid system has been described by Roger Brent and his colleagues (Gyuris, J. *et al.*, *Cell* 75:791-803 (1993); Zervos, A.S. *et al.*, *Cell* 72:223-232 (1993)).

By a "TNF-family ligand" is intended naturally occurring, recombinant, and synthetic ligands that are capable of binding to a member of the TNF receptor family and inducing the ligand/receptor signaling pathway. Members of the TNF ligand family include, but are not limited to, the TNFR-5, -6 $\alpha$  & -6 $\beta$  ligands, TNF- $\alpha$ , lymphdotoxin- $\alpha$  (LT- $\alpha$ , also known as TNF- $\beta$ ), LT- $\beta$ , FasL, CD40, CD27, CD30, 4-1BB, OX40 and nerve growth factor (NGF).

Representative therapeutic applications of the present invention are discussed in more detail below. The state of immunodeficiency that defines AIDS is secondary to a decrease in the number and function of CD4<sup>+</sup> T-lymphocytes. Recent reports estimate the daily loss of CD4<sup>+</sup> T cells to be between 3.5 X 10<sup>7</sup> and 2 X 10<sup>9</sup> cells (Wei X., *et al.*, *Nature* 373:117-122 (1995)). One cause of CD4<sup>+</sup> T cell depletion in the setting of HIV infection is believed to be HIV-induced apoptosis. Indeed, HIV-induced apoptotic cell death has been demonstrated not only *in vitro* but also, more importantly, in infected individuals (Ameisen, J.C., *AIDS* 8:1197-1213 (1994); Finkel, T.H., and Banda, N.K., *Curr. Opin. Immunol.* 6:605-615(1995); Muro-Cacho, C.A. *et al.*, *J. Immunol.* 154:5555-5566 (1995)). Furthermore, apoptosis and CD4<sup>+</sup> T-lymphocyte depletion is tightly correlated in different animal models of

AIDS (Brunner, T., *et al.*, *Nature* 373:441-444 (1995); Gougeon, M.L., *et al.*, *AIDS Res. Hum. Retroviruses* 9:553-563 (1993)) and, apoptosis is not observed in those animal models in which viral replication does not result in AIDS (Gougeon, M.L. *et al.*, *AIDS Res. Hum. Retroviruses* 9:553-563 (1993)). Further data indicates that uninfected but primed or activated T lymphocytes from HIV-infected individuals undergo apoptosis after encountering the TNF-family ligand FasL. Using monocytic cell lines that result in death following HIV infection, it has been demonstrated that infection of U937 cells with HIV results in the *de novo* expression of FasL and that FasL mediates HIV-induced apoptosis (Badley, A.D. *et al.*, *J. Virol.* 70:199-206 (1996)). Further the TNF-family ligand was detectable in uninfected macrophages and its expression was upregulated following HIV infection resulting in selective killing of uninfected CD4 T-lymphocytes (Badley, A.D. *et al.*, *J. Virol.* 70:199-206 (1996)). Thus, by the invention, a method for treating HIV<sup>+</sup> individuals is provided which involves administering an antagonist of the present invention to reduce selective killing of CD4 T-lymphocytes. Modes of administration and dosages are discussed in detail below.

In rejection of an allograft, the immune system of the recipient animal has not previously been primed to respond because the immune system for the most part is only primed by environmental antigens. Tissues from other members of the same species have not been presented in the same way that, for example, viruses and bacteria have been presented. In the case of allograft rejection, immunosuppressive regimens are designed to prevent the immune system from reaching the effector stage. However, the immune profile of xenograft rejection may resemble disease recurrence more than allograft rejection. In the case of disease recurrence, the immune system has already been activated, as evidenced by destruction of the native islet cells. Therefore, in disease recurrence the immune system is already at the effector stage. Agonist of the present invention are able to suppress the immune response to both allografts and xenografts because lymphocytes activated and differentiated into effector cells will express the TNFR polypeptide, and thereby are susceptible to compounds which enhance TNFR activity. Thus, the present invention further provides a method for creating immune privileged tissues. Antagonist of the invention can further be used in the treatment of Inflammatory Bowel-Disease.

### **Formulations**

The TNFR polypeptide composition will be formulated and dosed in a fashion consistent with good medical practice, taking into account the clinical condition of the individual patient (especially the side effects of treatment with TNFR-5, -6 $\alpha$  or -6 $\beta$  polypeptide alone), the site of delivery of the TNFR polypeptide composition, the method of administration, the scheduling of administration, and other factors known to practitioners. The "effective amount" of TNFR polypeptide for purposes herein is thus determined by such considerations.

As a general proposition, the total pharmaceutically effective amount of TNFR polypeptide administered parenterally per dose will be in the range of about 1  $\mu$ g/kg/day to 10 mg/kg/day of patient body weight, although, as noted above, this will be subject to therapeutic discretion. More preferably, this dose is at least 0.01 mg/kg/day, and most preferably for humans between about 0.01 and 1 mg/kg/day for the hormone. If given continuously, the TNFR polypeptide is typically administered at a dose rate of about 1  $\mu$ g/kg/hour to about 50  $\mu$ g/kg/hour, either by 1-4 injections per day or by continuous subcutaneous infusions, for example, using a mini-pump. An intravenous bag solution may also be employed. The length of treatment needed to observe changes and the interval following treatment for responses to occur appears to vary depending on the desired effect.

Pharmaceutical compositions containing the TNFR of the invention may be administered orally, rectally, parenterally, intracisternally, intravaginally, intraperitoneally, topically (as by powders, ointments, drops or transdermal patch), buccally, or as an oral or nasal spray. By "pharmaceutically acceptable carrier" is meant a non-toxic solid, semisolid or liquid filler, diluent, encapsulating material or formulation auxiliary of any type. The term "parenteral" as used herein refers to modes of administration



which include intravenous, intramuscular, intraperitoneal, intrasternal, subcutaneous and intraarticular injection and infusion.

The TNFR polypeptide is also suitably administered by sustained-release systems. Suitable examples of sustained-release compositions include semi-permeable polymer matrices in the form of shaped articles, e.g., films, or microcapsules. Sustained-release matrices include polylactides (U.S. Pat. No. 3,773,919, EP 58,481), copolymers of L-glutamic acid and gamma-ethyl-L-glutamate (Sidman, U. et al., *Biopolymers* 22:547-556 (1983)), poly (2-hydroxyethyl methacrylate) (R. Langer et al., *J. Biomed. Mater. Res.* 15:167-277 (1981), and R. Langer, *Chem. Tech.* 12:98-105 (1982)), ethylene vinyl acetate (R. Langer et al., *Id.*) or poly-D- (-)-3-hydroxybutyric acid (EP 133,988). Sustained-release TNFR polypeptide compositions also include liposomally entrapped TNFR polypeptides. Liposomes containing TNFR polypeptides are prepared by methods known per se: DE 3,218,121; Epstein et al., *Proc. Natl. Acad. Sci. (USA)* 82:3688-3692 (1985); Hwang et al., *Proc. Natl. Acad. Sci. (USA)* 77:4030-4034 (1980); EP 52,322; EP 36,676; EP 88,046; EP 143,949; EP 142,641; Japanese Pat. Appl. 83-118008; U.S. Pat. Nos. 4,485,045 and 4,544,545; and EP 102,324. Ordinarily, the liposomes are of the small (about 200-800 Angstroms) unilamellar type in which the lipid content is greater than about 30 mol. percent cholesterol, the selected proportion being adjusted for the optimal TNFR polypeptide therapy.

For parenteral administration, in one embodiment, the TNFR polypeptide is formulated generally by mixing it at the desired degree of purity, in a unit dosage injectable form (solution, suspension, or emulsion), with a pharmaceutically acceptable carrier, i.e., one that is non-toxic to recipients at the dosages and concentrations employed and is compatible with other ingredients of the formulation. For example, the formulation preferably does not include oxidizing agents and other compounds that are known to be deleterious to polypeptides.

Generally, the formulations are prepared by contacting the TNFR polypeptide uniformly and intimately with liquid carriers or finely divided solid carriers or both. Then, if necessary, the product is shaped into the desired formulation. Preferably the carrier is a parenteral carrier, more preferably a solution that is isotonic with the blood of the recipient. Examples of such carrier vehicles include water, saline, Ringer's solution, and dextrose solution. Non-aqueous vehicles such as fixed oils and ethyl oleate are also useful herein, as well as liposomes.

The carrier suitably contains minor amounts of additives such as substances that enhance isotonicity and chemical stability. Such materials are non-toxic to recipients at the dosages and concentrations employed, and include buffers such as phosphate, citrate, succinate, acetic acid, and other organic acids or their salts; antioxidants such as ascorbic acid; low molecular weight (less than about ten residues) polypeptides, e.g., polyarginine or tripeptides; proteins, such as serum albumin, gelatin, or immunoglobulins; hydrophilic polymers such as polyvinylpyrrolidone; amino acids, such as glycine, glutamic acid, aspartic acid, or arginine; monosaccharides, disaccharides, and other carbohydrates including cellulose or its derivatives, glucose, manose, or dextrans; chelating agents such as EDTA; sugar alcohols such as mannitol or sorbitol; counterions such as sodium; and/or nonionic surfactants such as polysorbates, poloxamers, or PEG.

The TNFR polypeptide is typically formulated in such vehicles at a concentration of about 0.1 mg/ml to 100 mg/ml, preferably 1-10 mg/ml, at a pH of about 3 to 8. It will be understood that the use of certain of the foregoing excipients, carriers, or stabilizers will result in the formation of TNFR polypeptide salts.

TNFR polypeptides to be used for therapeutic administration must be sterile. Sterility is readily accomplished by filtration through sterile filtration membranes (e.g., 0.2 micron membranes). Therapeutic TNFR polypeptide compositions generally are placed into a container having a sterile access port.

for example, an intravenous solution bag or vial having a stopper pierceable by a hypodermic injection needle.

TNFR polypeptides ordinarily will be stored in unit or multi-dose containers, for example, sealed ampoules or vials, as an aqueous solution or as a lyophilized formulation for reconstitution. As an example of a lyophilized formulation, 10-ml vials are filled with 5 ml of sterile-filtered 1% (w/v) aqueous TNFR polypeptide solution, and the resulting mixture is lyophilized. The infusion solution is prepared by reconstituting the lyophilized TNFR polypeptide using bacteriostatic Water-for-Injection.

The invention also provides a pharmaceutical pack or kit comprising one or more containers filled with one or more of the ingredients of the pharmaceutical compositions of the invention. Associated with such container(s) can be a notice in the form prescribed by a governmental agency regulating the manufacture, use or sale of pharmaceuticals or biological products, which notice reflects approval by the agency of manufacture, use or sale for human administration. In addition, the polypeptides of the present invention may be employed in conjunction with other therapeutic compounds.

#### *Chromosome Assays*

The nucleic acid molecules of the present invention are also valuable for chromosome identification. The sequence is specifically targeted to and can hybridize with a particular location on an individual human chromosome. Moreover, there is a current need for identifying particular sites on the chromosome. Few chromosome marking reagents based on actual sequence data (repeat polymorphisms) are presently available for marking chromosomal location. The mapping of DNAs to chromosomes according to the present invention is an important first step in correlating those sequences with genes associated with disease.

In certain preferred embodiments in this regard, the cDNAs herein disclosed are used to clone genomic DNA of a TNFR protein gene. This can

be accomplished using a variety of well known techniques and libraries, which generally are available commercially. The genomic DNA then is used for *in situ* chromosome mapping using well known techniques for this purpose.

In addition, in some cases, sequences can be mapped to chromosomes by preparing PCR primers (preferably 15-25 bp) from the cDNA. Computer analysis of the 3' untranslated region of the gene is used to rapidly select primers that do not span more than one exon in the genomic DNA, thus complicating the amplification process. These primers are then used for PCR screening of somatic cell hybrids containing individual human chromosomes. Fluorescence *in situ* hybridization ("FISH") of a cDNA clone to a metaphase chromosomal spread can be used to provide a precise chromosomal location in one step. This technique can be used with probes from the cDNA as short as 50 or 60 bp. For a review of this technique, see Verma *et al.*, *Human Chromosomes: A Manual Of Basic Techniques*, Pergamon Press, New York (1988).

Once a sequence has been mapped to a precise chromosomal location, the physical position of the sequence on the chromosome can be correlated with genetic map data. Such data are found, for example, in V. McKusick, *Mendelian Inheritance In Man*, available on-line through Johns Hopkins University, Welch Medical Library. The relationship between genes and diseases that have been mapped to the same chromosomal region are then identified through linkage analysis (coinheritance of physically adjacent genes).

Next, it is necessary to determine the differences in the cDNA or genomic sequence between affected and unaffected individuals. If a mutation is observed in some or all of the affected individuals but not in any normal individuals, then the mutation is likely to be the causative agent of the disease.

Having generally described the invention, the same will be more readily understood by reference to the following examples, which are provided by way of illustration and are not intended as limiting.

### Examples

#### *Example 1(a): Expression and Purification of "His-tagged" Extracellular form of TNFR-5 in E. coli*

The bacterial expression vector pQE9 (pD10) is used for bacterial expression in this example. (QIAGEN, Inc., 9259 Eton Avenue, Chatsworth, CA, 91311). pQE9 encodes ampicillin antibiotic resistance ("Amp<sup>r</sup>") and contains a bacterial origin of replication ("ori"), an IPTG inducible promoter, a ribosome binding site ("RBS"), six codons encoding histidine residues that allow affinity purification using nickel-nitrilo-tri-acetic acid ("Ni-NTA") affinity resin sold by QIAGEN, Inc., *supra*, and suitable single restriction enzyme cleavage sites. These elements are arranged such that an inserted DNA fragment encoding a polypeptide expresses that polypeptide with the six His residues (i.e., a "6 X His tag") covalently linked to the amino terminus of that polypeptide.

The DNA sequence encoding the desired portion of the TNFR-5 protein comprising the extracellular form of the TNFR-5 amino acid sequence is amplified from the deposited cDNA clone using PCR oligonucleotide primers which anneal to sequence encoding the amino terminal sequences of the desired portion of the TNFR-5 protein and to carboxy terminal sequences of the desired portion of the extracellular form of the TNFR-5 protein in the deposited cDNA. Additional nucleotides containing restriction sites to facilitate cloning in the pQE9 vector are added to the 5' and 3' primer sequences, respectively.

For cloning the extracellular form of the TNFR-5 protein, the 5' primer has the sequence 5' CGCGGATCCACCACTGCCCCGGCAGGAG 3' (SEQ ID NO: 25) containing the underlined BamHI restriction site followed by 18 nucleotides of the amino terminal coding sequence of the extracellular TNFR-5 sequence in SEQ ID NO:2. One of ordinary skill in the art would appreciate, of course, that the point in the protein coding sequence where the 5' primer

begins and where the 3' primer ends may be varied to amplify a DNA segment encoding any desired portion of the complete TNFR-5 protein shorter or longer than the extracellular form of the protein. The 3' primer has the sequence 5' GCGTCTAGACTAGTAATGAGAAGAGGCAGG 3' (SEQ ID NO:26) containing the underlined XbaI restriction site followed by 18 nucleotides complementary to the 3' end of cDNA encoding the extracellular domain of the TNFR-5 protein in Figure 1.

The amplified TNFR-5 DNA fragment and the vector pQE9 are digested with BamHI and XbaI and the digested DNAs are then ligated together. Insertion of the TNFR-5 DNA into the restricted pQE9 vector places the TNFR-5 protein coding region downstream from the IPTG-inducible promoter and in-frame with an initiating AUG and the six histidine codons.

The ligation mixture is transformed into competent *E. coli* cells using standard procedures such as those described in Sambrook et al., *Molecular Cloning: a Laboratory Manual, 2nd Ed.*; Cold Spring Harbor Laboratory Press, Cold Spring Harbor, NY (1989). *E. coli* strain M15/rep4, containing multiple copies of the plasmid pREP4, which expresses the lac repressor and confers kanamycin resistance ("Kan<sup>r</sup>"), is used in carrying out the illustrative example described herein. This strain, which is only one of many that are suitable for expressing TNFR-5 protein, is available commercially from QIAGEN, Inc., *supra*. Transformants are identified by their ability to grow on LB plates in the presence of ampicillin and kanamycin. Plasmid DNA is isolated from resistant colonies and the identity of the cloned DNA confirmed by restriction analysis, PCR and DNA sequencing.

Clones containing the desired constructs are grown overnight ("O/N") in liquid culture in LB media supplemented with both ampicillin (100 µg/ml) and kanamycin (25 µg/ml). The O/N culture is used to inoculate a large culture, at a dilution of approximately 1:25 to 1:250. The cells are grown to an

optical density at 600 nm ("OD600") of between 0.4 and 0.6. Isopropyl- $\beta$ -D-thiogalactopyranoside ("IPTG") is then added to a final concentration of 1 mM to induce transcription from the lac repressor sensitive promoter, by inactivating the lacI repressor. Cells subsequently are incubated further for 3 to 4 hours. Cells then are harvested by centrifugation.

The cells are then stirred for 3-4 hours at 4° C in 6M guanidine-HCl, pH 8. The cell debris is removed by centrifugation, and the supernatant containing the TNFR-5 is loaded onto a nickel-nitrilo-tri-acetic acid ("Ni-NTA") affinity resin column (available from QIAGEN, Inc., *supra*). Proteins with a 6 x His tag bind to the Ni-NTA resin with high affinity and can be purified in a simple one-step procedure (for details see: The QIAexpressionist, 1995, QIAGEN, Inc., *supra*). Briefly the supernatant is loaded onto the column in 6 M guanidine-HCl, pH 8, the column is first washed with 10 volumes of 6 M guanidine-HCl, pH 8, then washed with 10 volumes of 6 M guanidine-HCl pH 6, and finally the TNFR-5 is eluted with 6 M guanidine-HCl, pH 5.

The purified protein is then renatured by dialyzing it against phosphate-buffered saline (PBS) or 50 mM Na-acetate, pH 6 buffer plus 200 mM NaCl. Alternatively, the protein can be successfully refolded while immobilized on the Ni-NTA column. The recommended conditions are as follows: renature using a linear 6M-1M urea gradient in 500 mM NaCl, 20% glycerol, 20 mM Tris/HCl pH 7.4, containing protease inhibitors. The renaturation should be performed over a period of 1.5 hours or more. After renaturation the proteins can be eluted by the addition of 250 mM imidazole. Imidazole is removed by a final dialyzing step against PBS or 50 mM sodium acetate pH 6 buffer plus 200 mM NaCl. The purified protein is stored at 4° C or frozen at -80° C.

**Example 1(b): Expression and Purification of TNFR-6 $\alpha$  and -6 $\beta$  in *E. coli***

The bacterial expression vector pQE60 is used for bacterial expression in this example (QIAGEN, Inc., 9259 Eton Avenue, Chatsworth, CA, 91311). pQE60 encodes ampicillin antibiotic resistance ("Ampr") and contains a bacterial origin of replication ("ori"), an IPTG inducible promoter, a ribosome binding site ("RBS"), six codons encoding histidine residues that allow affinity purification using nickel-nitrilo-tri-acetic acid ("Ni-NTA") affinity resin sold by QIAGEN, Inc., *supra*, and suitable single restriction enzyme cleavage sites. These elements are arranged such that a DNA fragment encoding a polypeptide may be inserted in such a way as to produce that polypeptide with the six His residues (i.e., a "6 X His tag") covalently linked to the carboxyl terminus of that polypeptide. However, in this example, the polypeptide coding sequence is inserted such that translation of the six His codons is prevented and, therefore, the polypeptide is produced with no 6 X His tag.

The DNA sequences encoding the desired portions of TNFR-6 $\alpha$  and -6 $\beta$  proteins comprising the mature forms of the TNFR-6 $\alpha$  and -6 $\beta$  amino acid sequences are amplified from the deposited cDNA clones using PCR oligonucleotide primers which anneal to the amino terminal sequences of the desired portions of the TNFR-5, -6 $\alpha$  or -6 $\beta$  proteins and to sequences in the deposited constructs 3' to the cDNA coding sequence. Additional nucleotides containing restriction sites to facilitate cloning in the pQE60 vector are added to the 5' and 3' sequences, respectively.

For cloning the mature form of the TNFR-6 $\alpha$  protein, the 5' primer has the sequence 5' CGCCCATGGCCAGAAACACCCACCTAC 3' (SEQ ID NO:27) containing the underlined NcoI restriction site. One of ordinary skill in the art would appreciate, of course, that the point in the protein coding sequence where the 5' primer begins may be varied to amplify a desired



portion of the complete protein shorter or longer than the mature form. The 3' primer has the sequence 5' CGCAAGCTTCTCTTTTCAGTGCAAGTG 3' (SEQ ID NO:28) containing the underlined HindIII restriction site. For cloning the mature form of the TNFR-6 $\beta$  protein, the 5' primer has the sequence of SEQ ID NO:27 above, and the 3' primer has the sequence 5' CGCAAGCTTCTCTCAGCTCCTGCAGTG 3' (SEQ ID NO:29) containing the underlined HindIII restriction site.

The amplified TNFR-6 $\alpha$  and -6 $\beta$  DNA fragments and the vector pQE60 are digested with NcoI and HindIII and the digested DNAs are then ligated together. Insertion of the TNFR-6 $\alpha$  and -6 $\beta$  DNA into the restricted pQE60 vector places the TNFR-6 $\alpha$  and -6 $\beta$  protein coding region including its associated stop codon downstream from the IPTG-inducible promoter and in-frame with an initiating AUG. The associated stop codon prevents translation of the six histidine codons downstream of the insertion point.

The ligation mixture is transformed into competent *E. coli* cells using standard procedures such as those described in Sambrook et al., *Molecular Cloning: a Laboratory Manual, 2nd Ed.*; Cold Spring Harbor Laboratory Press, Cold Spring Harbor, NY (1989). *E. coli* strain M15/rep4, containing multiple copies of the plasmid pREP4, which expresses the lac repressor and confers kanamycin resistance ("Kanr"), is used in carrying out the illustrative example described herein. This strain, which is only one of many that are suitable for expressing TNFR-6 $\alpha$  or -6 $\beta$  protein, is available commercially from QIAGEN, Inc., *supra*. Transformants are identified by their ability to grow on LB plates in the presence of ampicillin and kanamycin. Plasmid DNA is isolated from resistant colonies and the identity of the cloned DNA confirmed by restriction analysis, PCR and DNA sequencing.

Clones containing the desired constructs are grown overnight ("O N") in liquid culture in LB media supplemented with both ampicillin (100  $\mu$ g/ml)

and kanamycin (25 µg/ml). The O/N culture is used to inoculate a large culture, at a dilution of approximately 1:25 to 1:250. The cells are grown to an optical density at 600 nm ("OD600") of between 0.4 and 0.6. isopropyl-b-D-thiogalactopyranoside ("IPTG") is then added to a final concentration of 1 mM to induce transcription from the *lac* repressor sensitive promoter, by inactivating the *lacI* repressor. Cells subsequently are incubated further for 3 to 4 hours. Cells then are harvested by centrifugation.

To purify the TNFR-6α and -6β polypeptide, the cells are then stirred for 3-4 hours at 4° C in 6M guanidine-HCl, pH 8. The cell debris is removed by centrifugation, and the supernatant containing the TNFR-6α and -6β is dialyzed against 50 mM Na-acetate buffer pH 6, supplemented with 200 mM NaCl. Alternatively, the protein can be successfully refolded by dialyzing it against 500 mM NaCl, 20% glycerol, 25 mM Tris/HCl pH 7.4, containing protease inhibitors. After renaturation the protein can be purified by ion exchange, hydrophobic interaction and size exclusion chromatography. Alternatively, an affinity chromatography step such as an antibody column can be used to obtain pure TNFR-6α and -6β protein. The purified protein is stored at 4° C or frozen at -80° C.

The following alternative method may be used to purify TNFR-5, -6α or -6β expressed in *E. coli* when it is present in the form of inclusion bodies. Unless otherwise specified, all of the following steps are conducted at 4-10°C.

Upon completion of the production phase of the *E. coli* fermentation, the cell culture is cooled to 4-10°C and the cells are harvested by continuous centrifugation at 15,000 rpm (Heraeus Sepatech). On the basis of the expected yield of protein per unit weight of cell paste and the amount of purified protein required, an appropriate amount of cell paste, by weight, is suspended in a buffer solution containing 100 mM Tris, 50 mM EDTA, pH 7.4. The cells are dispersed to a homogeneous suspension using a high shear mixer.

The cells were then lysed by passing the solution through a microfluidizer (Microfluidics, Corp. or APV Gaulin, Inc.) twice at 4000-6000 psi. The homogenate is then mixed with NaCl solution to a final concentration of 0.5 M NaCl, followed by centrifugation at 7000 xg for 15 min. The resultant pellet is washed again using 0.5M NaCl, 100 mM Tris, 50 mM EDTA, pH 7.4.

The resulting washed inclusion bodies are solubilized with 1.5 M guanidine hydrochloride (GuHCl) for 2-4 hours. After 7000 xg centrifugation for 15 min., the pellet is discarded and the TNFR-5, -6 $\alpha$  or -6 $\beta$  polypeptide-containing supernatant is incubated at 4°C overnight to allow further GuHCl extraction.

Following high speed centrifugation (30,000 xg) to remove insoluble particles, the GuHCl solubilized protein is refolded by quickly mixing the GuHCl extract with 20 volumes of buffer containing 50 mM sodium, pH 4.5, 150 mM NaCl, 2 mM EDTA by vigorous stirring. The refolded diluted protein solution is kept at 4°C without mixing for 12 hours prior to further purification steps.

To clarify the refolded TNF receptor polypeptide solution, a previously prepared tangential filtration unit equipped with 0.16  $\mu$ m membrane filter with appropriate surface area (e.g., Filtron), equilibrated with 40 mM sodium acetate, pH 6.0 is employed. The filtered sample is loaded onto a cation exchange resin (e.g., Poros HS-50, Perseptive Biosystems). The column is washed with 40 mM sodium acetate, pH 6.0 and eluted with 250 mM, 500 mM, 1000 mM, and 1500 mM NaCl in the same buffer, in a stepwise manner. The absorbance at 280 nm of the effluent is continuously monitored. Fractions are collected and further analyzed by SDS-PAGE.

Fractions containing the TNF receptor polypeptide are then pooled and mixed with 4 volumes of water. The diluted sample is then loaded onto a previously prepared set of tandem columns of strong anion (Poros HQ-50,

Perseptive Biosystems) and weak anion (Poros CM-20, Perseptive Biosystems) exchange resins. The columns are equilibrated with 40 mM sodium acetate, pH 6.0. Both columns are washed with 40 mM sodium acetate, pH 6.0, 200 mM NaCl. The CM-20 column is then eluted using a 10 column volume linear gradient ranging from 0.2 M NaCl, 50 mM sodium acetate, pH 6.0 to 1.0 M NaCl, 50 mM sodium acetate, pH 6.5. Fractions are collected under constant  $A_{280}$  monitoring of the effluent. Fractions containing the TNFR-5, -6 $\alpha$  or -6 $\beta$  polypeptide (determined, for instance, by 16% SDS-PAGE) are then pooled.

The resultant TNF receptor polypeptide exhibits greater than 95% purity after the above refolding and purification steps. No major contaminant bands are observed from Commassie blue stained 16% SDS-PAGE gel when 5  $\mu$ g of purified protein is loaded. The purified protein is also tested for endotoxin/LPS contamination, and typically the LPS content is less than 0.1 ng/ml according to LAL assays.

***Example 2: Cloning and Expression of TNFR-5, -6 $\alpha$  and -6 $\beta$  proteins in a Baculovirus Expression System***

In this illustrative example, the plasmid shuttle vector pA2 is used to insert the cloned DNA encoding complete protein, including its naturally associated secretory signal (leader) sequence, into a baculovirus to express the mature TNFR-5, -6 $\alpha$  or -6 $\beta$  protein, using standard methods as described in Summers *et al.*, *A Manual of Methods for Baculovirus Vectors and Insect Cell Culture Procedures*, Texas Agricultural Experimental Station Bulletin No. 1555 (1987). This expression vector contains the strong polyhedrin promoter of the *Autographa californica* nuclear polyhedrosis virus (AcMNPV) followed by convenient restriction sites such as BamHI, Xba I and Asp718. The polyadenylation site of the simian virus 40 ("SV40") is used for efficient polyadenylation. For easy selection of recombinant virus, the plasmid

contains the beta-galactosidase gene from *E. coli* under control of a weak *Drosophila* promoter in the same orientation, followed by the polyadenylation signal of the polyhedrin gene. The inserted genes are flanked on both sides by viral sequences for cell-mediated homologous recombination with wild-type viral DNA to generate a viable virus that express the cloned polynucleotide.

Many other baculovirus vectors could be used in place of the vector above, such as pAc373, pVL941 and pAcIM1, as one skilled in the art would readily appreciate, as long as the construct provides appropriately located signals for transcription, translation, secretion and the like, including a signal peptide and an in-frame AUG as required. Such vectors are described, for instance, in Luckow et al., *Virology* 170:31-39 (1989).

The cDNA sequence encoding the full length TNFR-5, -6 $\alpha$  or -6 $\beta$  protein in a deposited clone, including the AUG initiation codon and the naturally associated leader sequence shown in SEQ ID NO:2, 4, or 6 is amplified using PCR oligonucleotide primers corresponding to the 5' and 3' sequences of the gene. The 5' primer for TNFR-5 has the sequence 5' CGCTCTAGACCGCCATCATGGCCCGGATCCCCAAG 3' (SEQ ID NO:30) containing the underlined XbaI restriction enzyme site. The 5' primer for TNFR-6 $\alpha$  and -6 $\beta$  has the sequence

5' CGCGGATCCGCCATCATGAGGGCGTGGAGGGGCCAG 3' (SEQ ID NO:31) containing the underlined BamHI restriction enzyme site. All of the previously described primers encode an efficient signal for initiation of translation in eukaryotic cells, as described by Kozak, M., *J. Mol. Biol.* 196:947-950 (1987). The 3' primer for TNFR-5 has the sequence

5' GCGTCTAGACTAGTAATGAGAAGAGGCAGG 3' (SEQ ID NO:32) containing the underlined XbaI restriction site. The 3' primer for TNFR-6 $\alpha$  has the sequence 5' CGCCGTACCCTCTTTCAGTGCAAGTG 3' (SEQ ID NO:33) containing the underlined Asp718 restriction site. The 3' primer for

TNFR-6 $\beta$  has the sequence

5' CGCGGTACCCTCCTCAGCTCCTGCAGTG 3' (SEQ ID NO:34)

containing the underlined Asp718 restriction site.

The amplified fragment is isolated from a 1% agarose gel using a commercially available kit ("GeneClean," BIO 101 Inc., La Jolla, Ca.). The fragment then is digested with the appropriate restriction enzyme for each of the primers used, as specified above, and again is purified on a 1% agarose gel.

The plasmid is digested with the same restriction enzymes and optionally, can be dephosphorylated using calf intestinal phosphatase, using routine procedures known in the art. The DNA is then isolated from a 1% agarose gel using a commercially available kit ("GeneClean" BIO 101 Inc., La Jolla, Ca.).

The fragment and dephosphorylated plasmid are ligated together with T4 DNA ligase. *E. coli* HB101 or other suitable *E. coli* hosts such as XL-1 Blue (Statagene Cloning Systems, La Jolla, CA) cells are transformed with the ligation mixture and spread on culture plates. Bacteria are identified that contain the plasmid with the human TNF receptor gene by digesting DNA from individual colonies using the enzymes used immediately above and then analyzing the digestion product by gel electrophoresis. The sequence of the cloned fragment is confirmed by DNA sequencing. This plasmid is designated herein pA2-TNFR-5, pA2-TNFR-6 $\alpha$  or pA2TNFR-6 $\beta$  (collectively pA2-TNFR).

Five  $\mu$ g of the plasmid pA2-TNFR is co-transfected with 1.0  $\mu$ g of a commercially available linearized baculovirus DNA ("BaculoGold™ baculovirus DNA", Pharmingen, San Diego, CA), using the lipofection method described by Felgner et al., *Proc. Natl. Acad. Sci. USA* 84: 7413-7417 (1987). One  $\mu$ g of BaculoGold™ virus DNA and 5  $\mu$ g of the plasmid pA2-TNFR are mixed in a sterile well of a microtiter plate containing 50  $\mu$ l of serum-free Grace's medium (Life Technologies Inc., Gaithersburg, MD). Afterwards, 10

5  $\mu$ l Lipofectin plus 90  $\mu$ l Grace's medium are added, mixed and incubated for 15 minutes at room temperature. Then the transfection mixture is added drop-wise to Sf9 insect cells (ATCC CRL 1711) seeded in a 35 mm tissue culture plate with 1 ml Grace's medium without serum. The plate is then incubated for 5 hours at 27° C. The transfection solution is then removed from the plate and 1 ml of Grace's insect medium supplemented with 10% fetal calf serum is added. Cultivation is then continued at 27° C for four days.

10 After four days the supernatant is collected and a plaque assay is performed, as described by Summers and Smith, *supra*. An agarose gel with "Blue Gal" (Life Technologies Inc., Gaithersburg) is used to allow easy identification and isolation of gal-expressing clones, which produce blue-stained plaques. (A detailed description of a "plaque assay" of this type can also be found in the user's guide for insect cell culture and baculovirology distributed by Life Technologies Inc., Gaithersburg, page 9-10). After appropriate incubation, blue stained plaques are picked with the tip of a micropipettor (e.g., Eppendorf). The agar containing the recombinant viruses is then resuspended in a microcentrifuge tube containing 200  $\mu$ l of Grace's medium and the suspension containing the recombinant baculovirus is used to infect Sf9 cells seeded in 35 mm dishes. Four days later the supernatants of these culture dishes are harvested and then they are stored at 4° C.

15 To verify the expression of the TNF receptor gene Sf9 cells are grown in Grace's medium supplemented with 10% heat-inactivated FBS. The cells are infected with the recombinant baculovirus at a multiplicity of infection ("MOI") of about 2. If radiolabeled proteins are desired, 6 hours later the medium is removed and is replaced with SF900 II medium minus methionine and cysteine (available from Life Technologies Inc., Rockville, MD). After 42 hours, 5  $\mu$ Ci of  $^{35}$ S-methionine and 5  $\mu$ Ci  $^{35}$ S-cysteine (available from Amersham) are added. The cells are further incubated for 16 hours and then are harvested by centrifugation. The proteins in the supernatant as well as the

intracellular proteins are analyzed by SDS-PAGE followed by autoradiography (if radiolabeled).

Microsequencing of the amino acid sequence of the amino terminus of purified protein may be used to determine the amino terminal sequence of the mature form of the TNF receptor protein.

*Example 3: Cloning and Expression of TNFR-5, -6 $\alpha$  and -6 $\beta$  in Mammalian Cells*

A typical mammalian expression vector contains the promoter element, which mediates the initiation of transcription of mRNA, the protein coding sequence, and signals required for the termination of transcription and polyadenylation of the transcript. Additional elements include enhancers, Kozak sequences and intervening sequences flanked by donor and acceptor sites for RNA splicing. Highly efficient transcription can be achieved with the early and late promoters from SV40, the long terminal repeats (LTRs) from Retroviruses, e.g., RSV, HTLV1, HIV1 and the early promoter of the cytomegalovirus (CMV). However, cellular elements can also be used (e.g., the human actin promoter). Suitable expression vectors for use in practicing the present invention include, for example, vectors such as pSVL and pMSG (Pharmacia, Uppsala, Sweden), pRSVcat (ATCC 37152), pSV2dhfr (ATCC 37146) and pBC12MI (ATCC 67109). Mammalian host cells that could be used include, human Hela, 293, H9 and Jurkat cells, mouse NIH3T3 and C127 cells, Cos 1, Cos 7 and CV1, quail QC1-3 cells, mouse L cells and Chinese hamster ovary (CHO) cells.

Alternatively, the gene can be expressed in stable cell lines that contain the gene integrated into a chromosome. The co-transfection with a selectable marker such as dhfr, gpt, neomycin, hygromycin allows the identification and isolation of the transfected cells.



The transfected gene can also be amplified to express large amounts of the encoded protein. The DHFR (dihydrofolate reductase) marker is useful to develop cell lines that carry several hundred or even several thousand copies of the gene of interest. Another useful selection marker is the enzyme glutamine synthase (GS) (Murphy et al., *Biochem J.* 227:277-279 (1991); Bebbington et al., *Bio/Technology* 10:169-175 (1992)). Using these markers, the mammalian cells are grown in selective medium and the cells with the highest resistance are selected. These cell lines contain the amplified gene(s) integrated into a chromosome. Chinese hamster ovary (CHO) and NSO cells are often used for the production of proteins.

The expression vectors pC1 and pC4 contain the strong promoter (LTR) of the Rous Sarcoma Virus (Cullen et al., *Molecular and Cellular Biology*, 438-447 (March, 1985)) plus a fragment of the CMV-enhancer (Boshart et al., *Cell* 41:521-530 (1985)). Multiple cloning sites, e.g., with the restriction enzyme cleavage sites BamHI, XbaI and Asp718, facilitate the cloning of the gene of interest. The vectors contain in addition the 3' intron, the polyadenylation and termination signal of the rat preproinsulin gene.

#### **Example 3(a): Cloning and Expression in COS Cells**

The expression plasmid, pTNFR-5-HA, is made by cloning a portion of the cDNA encoding the mature form of the TNF receptor protein into the expression vector pcDNA1 Amp or pcDNAIII (which can be obtained from Invitrogen, Inc.).

The expression vector pcDNA1 amp contains: (1) an *E. coli* origin of replication effective for propagation in *E. coli* and other prokaryotic cells; (2) an ampicillin resistance gene for selection of plasmid-containing prokaryotic cells; (3) an SV40 origin of replication for propagation in eukaryotic cells; (4) a CMV promoter, a polylinker, an SV40 intron; (5) several codons encoding a hemagglutinin fragment (i.e., an "HA" tag to facilitate purification) followed by

a termination codon and polyadenylation signal arranged so that a cDNA can be conveniently placed under expression control of the CMV promoter and operably linked to the SV40 intron and the polyadenylation signal by means of restriction sites in the polylinker. The HA tag corresponds to an epitope  
5 derived from the influenza hemagglutinin protein described by Wilson et al., *Cell* 37: 767 (1984). The fusion of the HA tag to the target protein allows easy detection and recovery of the recombinant protein with an antibody that recognizes the HA epitope. pcDNAIII contains, in addition, the selectable neomycin marker.

10 A DNA fragment encoding the complete TNF receptor polypeptide is cloned into the polylinker region of the vector so that recombinant protein expression is directed by the CMV promoter. The plasmid construction strategy is as follows. The TNF receptor cDNA of a deposited clone is amplified using primers that contain convenient restriction sites, much as  
15 described above for construction of vectors for expression of a TNF receptor in *E. coli*. Suitable primers include the following, which are used in this example. The 5' primer for TNFR-5, containing the underlined EcoRI site, has the following sequence:

5' CGCGAATTCCGCCATCATGGCCCGGATCCCCAAG 3' (SEQ ID

20 NO:35). The 3' primer, containing the underlined XbaI site, has the following sequence: 5' GCGTCTAGAGTAATGAGAAGAGGCAGG 3' (SEQ ID NO:36).

25 The PCR amplified DNA fragment and the vector, pcDNAI/Amp, are digested with XbaI and EcoRI and then ligated. The ligation mixture is transformed into *E. coli* strain SURE (available from Stratagene Cloning Systems, 11099 North Torrey Pines Road, La Jolla, CA 92037), and the transformed culture is plated on ampicillin media plates which then are incubated to allow growth of ampicillin resistant colonies. Plasmid DNA is

isolated from resistant colonies and examined by restriction analysis or other means for the presence of the fragment encoding the TNFR-5 polypeptide

For expression of recombinant TNFR-5, COS cells are transfected with an expression vector, as described above, using DEAE-DEXTRAN, as described, for instance, in Sambrook et al., *Molecular Cloning: a Laboratory Manual*, Cold Spring Laboratory Press, Cold Spring Harbor, New York (1989). Cells are incubated under conditions for expression of TNFR-5 by the vector.

Expression of the pTNFR-5-HA fusion protein is detected by radiolabeling and immunoprecipitation, using methods described in, for example Harlow et al., *Antibodies: A Laboratory Manual, 2nd Ed.*; Cold Spring Harbor Laboratory Press, Cold Spring Harbor, New York (1988). To this end, two days after transfection, the cells are labeled by incubation in media containing  $^{35}\text{S}$ -cysteine for 8 hours. The cells and the media are collected, and the cells are washed and the lysed with detergent-containing RIPA buffer: 150 mM NaCl, 1% NP-40, 0.1% SDS, 1% NP-40, 0.5% DOC, 50 mM TRIS, pH 7.5, as described by Wilson et al. cited above. Proteins are precipitated from the cell lysate and from the culture media using an HA-specific monoclonal antibody. The precipitated proteins then are analyzed by SDS-PAGE and autoradiography. An expression product of the expected size is seen in the cell lysate, which is not seen in negative controls.

#### *Example 3(b): Cloning and Expression in CHO Cells*

The vector pC4 is used for the expression of TNFR-5, -6 $\alpha$  and -6 $\beta$  polypeptides. Plasmid pC4 is a derivative of the plasmid pSV2-dhfr (ATCC Accession No. 37146). The plasmid contains the mouse DHFR gene under control of the SV40 early promoter. Chinese hamster ovary- or other cells lacking dihydrofolate activity that are transfected with these plasmids can be selected by growing the cells in a selective medium (alpha minus MEM, Life

Technologies) supplemented with the chemotherapeutic agent methotrexate. The amplification of the DHFR genes in cells resistant to methotrexate (MTX) has been well documented (see, e.g., Alt, F. W., Kellems, R. M., Bertino, J. R., and Schimke, R. T., 1978, *J. Biol. Chem.* 253:1357-1370, Hamlin, J. L. and Ma, C. 1990, *Biochem. et Biophys. Acta*, 1097:107-143, Page, M. J. and Sydenham, M. A. 1991, *Biotechnology* 9:64-68). Cells grown in increasing concentrations of MTX develop resistance to the drug by overproducing the target enzyme, DHFR, as a result of amplification of the DHFR gene. If a second gene is linked to the DHFR gene, it is usually co-amplified and over-expressed. It is known in the art that this approach may be used to develop cell lines carrying more than 1,000 copies of the amplified gene(s). Subsequently, when the methotrexate is withdrawn, cell lines are obtained which contain the amplified gene integrated into one or more chromosome(s) of the host cell.

Plasmid pC4 contains for expressing the gene of interest the strong promoter of the long terminal repeat (LTR) of the Rouse Sarcoma Virus (Cullen, et al., *Molecular and Cellular Biology*, March 1985:438-447) plus a fragment isolated from the enhancer of the immediate early gene of human cytomegalovirus (CMV) (Boshart et al., *Cell* 41:521-530 (1985)). Downstream of the promoter are the following single restriction enzyme cleavage sites that allow the integration of the genes: BamHI, Xba I, and Asp718. Behind these cloning sites the plasmid contains the 3' intron and polyadenylation site of the rat preproinsulin gene. Other high efficiency promoters can also be used for the expression, e.g., the human  $\beta$ -actin promoter, the SV40 early or late promoters or the long terminal repeats from other retroviruses, e.g., HIV and HTLV. Clontech's Tet-Off and Tet-On gene expression systems and similar systems can be used to express the TNF receptor polypeptide in a regulated way in mammalian cells (Gossen, M., & Bujard, H. 1992, *Proc. Natl. Acad. Sci. USA* 89:5547-5551). For the

polyadenylation of the mRNA other signals, e.g., from the human growth hormone or globin genes can be used as well. Stable cell lines carrying a gene of interest integrated into the chromosomes can also be selected upon co-transfection with a selectable marker such as gpt, G418 or hygromycin. It is advantageous to use more than one selectable marker in the beginning, e.g., G418 plus methotrexate.

The plasmid pC4 is digested with the restriction enzymes appropriate for the specific primers used to amplify the TNF receptor of choice as outlined below and then dephosphorylated using calf intestinal phosphates by procedures known in the art. The vector is then isolated from a 1% agarose gel.

The DNA sequence encoding the TNF receptor polypeptide is amplified using PCR oligonucleotide primers corresponding to the 5' and 3' sequences of the desired portion of the gene. The 5' primer for TNFR-5 containing the underlined XbaI site, has the following sequence:

5' CGCTCTAGACCGCCATCATGGCCCGGATCCCCAAG 3' (SEQ ID NO:37). The 5' primer for TNFR-6 $\alpha$  and -6 $\beta$  containing the underlined

BamHI site, has the following sequence:

5' CGCGGATCCCGCCATCATGAGGGCGTGGAGGGGCCAG 3' (SEQ ID NO:31). The 3' primer for TNFR-5, containing the underlined XbaI site,

has the following sequence:

5' GCGTCTAGACTAGTAATGAGAAGAGGCAGG 3' (SEQ ID NO:38).

The 3' primer for TNFR-6 $\alpha$  has the sequence

5' CGCGGTACCCTCTTTTCAGTGCAAGTG 3' (SEQ ID NO:33)

containing the underlined Asp718 restriction site. The 3' primer for TNFR-6 $\beta$

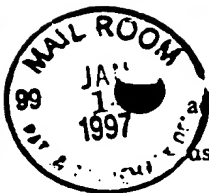
has the sequence 5' CGCGGTACCCTCCTCAGCTCCTGCAGTG 3' (SEQ ID NO:34) containing the underlined Asp718 restriction site.

The amplified fragment is digested with the endonucleases which will cut at the engineered restriction site(s) and then purified again on a 1% agarose gel. The isolated fragment and the dephosphorylated vector are then ligated with T4 DNA ligase. *E. coli* HB101 or XL-1 Blue cells are then transformed and bacteria are identified that contain the fragment inserted into plasmid pC4 using, for instance, restriction enzyme analysis.

Chinese hamster ovary cells lacking an active DHFR gene are used for transfection. Five  $\mu$ g of the expression plasmid pC4 is cotransfected with 0.5  $\mu$ g of the plasmid pSVneo using lipofectin (Felgner et al., *supra*). The plasmid pSV2-neo contains a dominant selectable marker, the *neo* gene from Tn5 encoding an enzyme that confers resistance to a group of antibiotics including G418. The cells are seeded in alpha minus MEM supplemented with 1 mg/ml G418. After 2 days, the cells are trypsinized and seeded in hybridoma cloning plates (Greiner, Germany) in alpha minus MEM supplemented with 10, 25, or 50 ng/ml of methotrexate plus 1 mg/ml G418. After about 10-14 days single clones are trypsinized and then seeded in 6-well petri dishes or 10 ml flasks using different concentrations of methotrexate (50 nM, 100 nM, 200 nM, 400 nM, 800 nM). Clones growing at the highest concentrations of methotrexate are then transferred to new 6-well plates containing even higher concentrations of methotrexate (1  $\mu$ M, 2  $\mu$ M, 5  $\mu$ M, 10 mM, 20 mM). The same procedure is repeated until clones are obtained which grow at a concentration of 100 - 200  $\mu$ M. Expression of the desired gene product is analyzed, for instance, by SDS-PAGE and Western blot or by reversed phase HPLC analysis.

**Example 4: Tissue distribution of TNF receptor mRNA expression**

Northern blot analysis is carried out to examine TNFR-5, -6 $\alpha$  or -6 $\beta$  gene expression in human tissues, using methods described by, among others, Sambrook *et al.*, cited above. A cDNA probe containing the entire nucleotide sequence of a TNF receptor protein (SEQ ID NO:1, 3 or 5) is labeled with  $^{32}$ P using the *rediprime*<sup>TM</sup> DNA labeling system (Amersham Life Science).

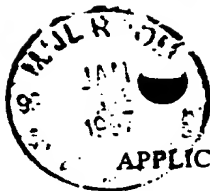


according to manufacturer's instructions. After labeling, the probe is purified using a CHROMA SPIN-100™ column (Clontech Laboratories, Inc.), according to manufacturer's protocol number PT1200-1. The purified labeled probe is then used to examine various human tissues for TNF receptor mRNA.

5 Multiple Tissue Northern (MTN) blots containing various human tissues (H) or human immune system tissues (IM) are obtained from Clontech and are examined with the labeled probe using ExpressHyb™ hybridization solution (Clontech) according to manufacturer's protocol number PT1190-1. Following hybridization and washing, the blots are mounted and exposed to  
10 film at -70° C overnight, and films developed according to standard procedures.

It will be clear that the invention may be practiced otherwise than as particularly described in the foregoing description and examples. Numerous modifications and variations of the present invention are possible in light of  
15 the above teachings and, therefore, are within the scope of the appended claims.

The entire disclosure of all publications (including patents, patent applications, journal articles, laboratory manuals, books, or other documents) cited herein are hereby incorporated by reference.



IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

APPLICANTS : WEI *et al.*,  
APPLICATION NO. : Unassigned  
FILED : Herewith  
FOR : Tumor Necrosis Factor Receptors 5, 6 Alpha and 6  
Beta :  
ATTORNEY DOCKET NO: PF341

STATEMENT UNDER 37 C.F.R. 1.821(a)

Assistant Commissioner For Patents  
Washington, D.C. 20231

January 15, 1997

Sir:

Applicants hereby certify that the hard copy of the sequence listing and the computer-readable form of such sequence listing are identical.

Respectfully submitted,

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WP PTO Forms Statmt. 1.821



## SEQUENCE LISTING

## (1) GENERAL INFORMATION:

- (i) APPLICANT: WEI, YING-FEI  
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GENTZ, REINER  
RUBEN, STEVEN
- (ii) TITLE OF INVENTION: TUMOR NECROSIS FACTOR RECEPTORS 5, 6  
ALPHA AND 6 BETA
- (iii) NUMBER OF SEQUENCES: 38
- (iv) CORRESPONDENCE ADDRESS:
  - (A) ADDRESSEE: HUMAN GENOME SCIENCES, INC.
  - (B) STREET: 9410 KEY WEST AVENUE
  - (C) CITY: ROCKVILLE
  - (D) STATE: MD
  - (E) COUNTRY: US
  - (F) ZIP: 20850
- (v) COMPUTER READABLE FORM:
  - (A) MEDIUM TYPE: Floppy disk
  - (B) COMPUTER: IBM PC compatible
  - (C) OPERATING SYSTEM: PC-DOS/MS-DOS
  - (D) SOFTWARE: PatentIn Release #1.0, Version #1.30
- (vi) CURRENT APPLICATION DATA:
  - (A) APPLICATION NUMBER:
  - (B) FILING DATE:
  - (C) CLASSIFICATION:
- (viii) ATTORNEY/AGENT INFORMATION:
  - (A) NAME: BROOKES, ANDERS A
  - (B) REGISTRATION NUMBER: 36,373
  - (C) REFERENCE/DOCKET NUMBER: PF341
- (ix) TELECOMMUNICATION INFORMATION:
  - (A) TELEPHONE: (301) 309-8504
  - (B) TELEFAX: (301) 309-8512

## (2) INFORMATION FOR SEQ ID NO:1:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 1392 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (genomic)

(ix) FEATURES:

(A) NAME/KEY: CDS

(B) LOCATION: 183..959

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:1:

CCTCTCCACG	GGACGAACT	CAGCCAACGA	TTTCTGATAG	ATTTTTGGGA	GTTTGACCAG	60
AGATGCAAGG	GGAGAGGAG	CGCTTCCTAC	CGTTAGGGAA	CTCTGGGGAC	AGAGCGCCCC	120
GGCCGCTGA	TGGCCGCGC	AGGGTGCGAC	CCAGGACCCA	GGACGGCGTC	GGGAACCATA	180
CC ATG GCC CGG ATC CCC AAG ACC CTA AAG TTC GTC GTC GTC ATC GTC						227
Met Ala Arg Ile Pro Lys Thr Leu Lys Phe Val Val Val Ile Val						
1	5		10		15	
GCG GTC CTG CTG CCA CTC CTA GCT TAC TCT GCC ACC ACT GCC CGG CAG						275
Ala Val Leu Leu Pro Val Leu Ala Tyr Ser Ala Thr Thr Ala Arg Gln						
	20		25		30	
GAG GAA GTT CCC CAG CAG ACA GTG GCC CCA CAG CAA CAG AGG CAC AGC						323
Glu Glu Val Pro Gln Gln Thr Val Ala Pro Gln Gln Gln Arg His Ser						
	35		40		45	
TTC AAG GGG GAG GAG TGT CCA GCA GGA TCT CAT AGA TCA GAA CAT ACT						371
Phe Lys Gly Glu Glu Cys Pro Ala Gly Ser His Arg Ser Glu His Thr						
	50		55		60	
GGA GCC TGT AAC CCG TGC ACA GAG GGT GTG GAT TAC ACC AAC GCT TCC						419
Gly Ala Cys Asn Pro Cys Thr Glu Gly Val Asp Tyr Thr Asn Ala Ser						
	65		70		75	
AAC AAT GAA CCT TCT TGC TTC CCA TGT ACA GTT TGT AAA TCA GAT CAA						467
Asn Asn Glu Pro Ser Cys Phe Pro Cys Thr Val Cys Lys Ser Asp Gln						
	80		85		90	95
AAA CAT AAA AGT TCC TGC ACC ATG ACC AGA CAC ACA GTG TGT CAG TGT						515
Lys His Lys Ser Ser Cys Thr Met Thr Arg Asp Thr Val Cys Gln Cys						
	100		105		110	
AAA GAA GGC ACC TTC CGG AAT GAA AAC TCC CCA GAG ATG TGC CGG AAG						563
Lys Glu Gly Thr Phe Arg Asn Glu Asn Ser Pro Glu Met Cys Arg Lys						
	115		120		125	
TGT AGC AGG TGC CCT AGT GGG GAA GTC CAA GTC AGT AAT TGT ACC TCC						611
Cys Ser Arg Cys Pro Ser Gly Glu Val Gln Val Ser Asn Cys Thr Ser						
	130		135		140	
TGG GAT GAT ATC CAG TGT GTT GAA GAA TTT GGT GCC AAT GCC ACT GTG						659
Trp Asp Asp Ile Gln Cys Val Glu Glu Phe Gly Ala Asn Ala Thr Val						
	145		150		155	

GAA ACC CCA GCT GCT GAA GAC ACA ATG AAC ACC AGC CCG GGG ACT CCT Glu Thr Pro Ala Ala Glu Glu Thr Met Asn Thr Ser Pro Gly Thr Pro 160 165 170 175	707
GCC CCA GCT GCT GAA GAG ACA ATG AAC ACC AGC CCA GGG ACT CCT GCC Ala Pro Ala Ala Glu Glu Thr Met Asn Thr Ser Pro Gly Thr Pro Ala 180 185 190	755
CCA GCT GCT GAA GAC ACA ATG ACC ACC AGC CCG GGG ACT CCT GCC CCA Pro Ala Ala Glu Glu Thr Met Thr Thr Ser Pro Gly Thr Pro Ala Pro 195 200 205	803
GCT GCT GAA GAG ACA ATG ACC ACC AGC CCG GGG ACT CCT GCC CCA GCT Ala Ala Glu Glu Thr Met Thr Thr Ser Pro Gly Thr Pro Ala Pro Ala 210 215 220	851
GCT GAA GAG ACA ATG ACC ACC AGC CCG GGG ACT CCT GCC TCT TCT CAT Ala Glu Glu Thr Met Thr Thr Ser Pro Gly Thr Pro Ala Ser Ser His 225 230 235	899
TAC CTC TCA TGC ACC ATC GTA GGG ATC ATA GTT CTA ATT GTG CTT CTG Tyr Leu Ser Cys Thr Ile Val Gly Ile Ile Val Leu Ile Val Leu Leu 240 245 250 255	947
ATT GTG TTT GTT TGAAAGACTT CACTGTGGAA GAAATTCCTT CCTTACCTGA Ile Val Phe Val	999
AAGGTTTCAGG TAGGCGCTGG CTGAGGGCGG GGGGCGCTGG ACACTCTCTG CCTGCTCTCC	1059
CTCTGCTGTG TTCCACACA CAGAAACGCC TGCCCTGCC CCAAGTCTG GTGTCTCCAG	1119
CCTGGCTCTA TCTTCTCTT TGTGATCGTC CCATCCCCAC ATCCCGTGCA CCCCCAGGA	1179
CCCTGGTCTC ATCAGTCCCT CTCCTGGAGC TGGGGGTCCA CACATCTCCC AGCCAAGTCC	1239
AAGAGGCAGG GCCAGTTCCT CCCATCTTCA GGCCAGCCA GGCAGGGGGC AGTCGCCTCC	1299
TCAACTGGGT GACAAGGGTG AGGATGAGAA GTGGTCACGG GATTATTCA GCCTTGGTCA	1359
GAGCAGAACA CAGAGATTTT CCGTGAAAAA AAA	1392

## (2) INFORMATION FOR SEQ ID NO:2:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 259 amino acids
- (B) TYPE: amino acid
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: protein

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:2:

Met Ala Arg Ile Pro Lys Thr Leu Lys Phe Val Val Val Ile Val Ala  
 1 5 10 15  
 Val Leu Leu Pro Val Leu Ala Tyr Ser Ala Thr Thr Ala Arg Gln Glu  
 20 25 30  
 Glu Val Pro Gln Gln Thr Val Ala Pro Gln Gln Gln Arg His Ser Phe  
 35 40 45  
 Lys Gly Glu Glu Cys Pro Ala Gly Ser His Arg Ser Glu His Thr Gly  
 50 55 60  
 Ala Cys Asn Pro Cys Thr Glu Gly Val Asp Tyr Thr Asn Ala Ser Asn  
 65 70 75 80  
 Asn Glu Pro Ser Cys Phe Pro Cys Thr Val Cys Lys Ser Asp Gln Lys  
 85 90 95  
 His Lys Ser Ser Cys Thr Met Thr Arg Asp Thr Val Cys Gln Cys Lys  
 100 105 110  
 Glu Gly Thr Phe Arg Asn Glu Asn Ser Pro Glu Met Cys Arg Lys Cys  
 115 120 125  
 Ser Arg Cys Pro Ser Gly Glu Val Gln Val Ser Asn Cys Thr Ser Trp  
 130 135 140  
 Asp Asp Ile Gln Cys Val Glu Glu Phe Gly Ala Asn Ala Thr Val Glu  
 145 150 155 160  
 Thr Pro Ala Ala Glu Glu Thr Met Asn Thr Ser Pro Gly Thr Pro Ala  
 165 170 175  
 Pro Ala Ala Glu Glu Thr Met Asn Thr Ser Pro Gly Thr Pro Ala Pro  
 180 185 190  
 Ala Ala Glu Glu Thr Met Thr Thr Ser Pro Gly Thr Pro Ala Pro Ala  
 195 200 205  
 Ala Glu Glu Thr Met Thr Thr Ser Pro Gly Thr Pro Ala Pro Ala Ala  
 210 215 220  
 Glu Glu Thr Met Thr Thr Ser Pro Gly Thr Pro Ala Ser Ser His Tyr  
 225 230 235 240  
 Leu Ser Cys Thr Ile Val Gly Ile Ile Val Leu Ile Val Leu Leu Ile  
 245 250 255  
 Val Phe Val

(2) INFORMATION FOR SEQ ID NO:3:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1077 base pairs

(B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(ix) FEATURE:

(A) NAME/KEY: CDS  
 (B) LOCATION: 25..924

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:3:

GCTCTCCCTG CTCCAGCAAG GACC ATG AGG GCG CTG GAG GGG CCA GGC CTG	51
Met Arg Ala Leu Glu Gly Pro Gly Leu	
1 5	
TCG CTG CTG TGC CTG GTG TTG GCG CTG CCT GCC CTG CTG CCG GTG CCG	99
Ser Leu Leu Cys Leu Val Leu Ala Leu Pro Ala Leu Leu Pro Val Pro	
10 15 20 25	
GCT GTA CGC GGA GTG GCA GAA ACA CCC ACC TAC CCC TGG CCG GAC GCA	147
Ala Val Arg Gly Val Ala Glu Thr Pro Thr Tyr Pro Trp Arg Asp Ala	
30 35 40	
GAG ACA GGG GAG CCG CTG GTG TGC GCC CAG TGC CCC CCA GGC ACC TTT	195
Glu Thr Gly Glu Arg Leu Val Cys Ala Gln Cys Pro Pro Gly Thr Phe	
45 50 55	
GTG CAG CCG CCG TGC CGC CGA GAC AGC CCC ACG ACG TGT GGC CCG TGT	243
Val Gln Arg Pro Cys Arg Arg Asp Ser Pro Thr Thr Cys Gly Pro Cys	
60 65 70	
CCA CCG CGC CAC TAC ACG CAG TTC TGG AAC TAC CTG GAG CGC TGC CGC	291
Pro Pro Arg His Tyr Thr Gln Phe Trp Asn Tyr Leu Glu Arg Cys Arg	
75 80 85	
TAC TGC AAC GTC CTC TGC GGG GAG CGT GAG GAG GAG GCA CCG GCT TGC	339
Tyr Cys Asn Val Leu Cys Gly Glu Arg Glu Glu Glu Ala Arg Ala Cys	
90 95 100 105	
CAC GCC ACC CAC AAC CGT GCC TGC CGC TGC CGC ACC GGC TTC TTC GCG	387
His Ala Thr His Asn Arg Ala Cys Arg Cys Arg Thr Gly Phe Phe Ala	
110 115 120	
CAC GCT GGT TTC TGC TTG GAG CAC GCA TCG TGT CCA CCT GGT GCC GGC	435
His Ala Gly Phe Cys Leu Glu His Ala Ser Cys Pro Pro Gly Ala Gly	
125 130 135	
GTG ATT GCC CCG GGC ACC CCC AGC CAG AAC ACG CAG TGC CAG CCG TGC	483
Val Ile Ala Pro Gly Thr Pro Ser Gln Asn Thr Gln Cys Gln Pro Cys	
140 145 150	

CCC CCA GGC ACC TTC TCA GCC AGC AGC TCC AGC TCA GAG CAG TGC CAG	531
Pro Pro Gly Thr Phe Ser Ala Ser Ser Ser Ser Ser Glu Gln Cys Gln	
155 160 165	
CCC CAC CGC AAC TGC ACG GCC CTG GGC CTG GCC CTC AAT GTG CCA GGC	579
Pro His Arg Asn Cys Thr Ala Leu Gly Leu Ala Leu Asn Val Pro Gly	
170 175 180 185	
TCT TCC TCC CAT GAC ACC CTG TGC ACC AGC TGC ACT GGC TTC CCC CTC	627
Ser Ser Ser His Asp Thr Leu Cys Thr Ser Cys Thr Gly Phe Pro Leu	
190 195 200	
AGC ACC AGG GTA CCA GGA GCT GAG GAG TGT GAG CGT GCC GTC ATC GAC	675
Ser Thr Arg Val Pro Gly Ala Glu Glu Cys Glu Arg Ala Val Ile Asp	
205 210 215	
TTT GTG GCT TTC CAG GAC ATC TCC ATC AAG AGG CTG CAG CGG CTG CTG	723
Phe Val Ala Phe Gln Asp Ile Ser Ile Lys Arg Leu Gln Arg Leu Leu	
220 225 230	
CAG GCC CTC GAG GCC CCG GAG GGC TGG GGT CCG ACA CCA AGG GCG GGC	771
Gln Ala Leu Glu Ala Pro Glu Gly Trp Gly Pro Thr Pro Arg Ala Gly	
235 240 245	
CGC GCG GCC TTG CAG CTG AAG CTG CGT CGG CGG CTC ACG GAG CTC CTG	819
Arg Ala Ala Leu Gln Leu Lys Leu Arg Arg Arg Leu Thr Glu Leu Leu	
250 255 260 265	
GGG GCG CAG GAC GGG GCG CTG CTG GTG CGG CTG CTG CAG GCG CTG CGC	867
Gly Ala Gln Asp Gly Ala Leu Leu Val Arg Leu Leu Gln Ala Leu Arg	
270 275 280	
GTG GCC AGG ATG CCC GGG CTG GAG CGG AGC GTC CGT GAG CGC TTC CTC	915
Val Ala Arg Met Pro Gly Leu Glu Arg Ser Val Arg Glu Arg Phe Leu	
285 290 295	
CCT GTG CAC TGATCCTGGC CCCCTCTTAT TTATTCTACA TCCTTGGCAC	964
Pro Val His	
300	
CCCACTTGCA CTGAAAGAGG CTTTTTTTAA AATAGAAGAA ATGAGGTTTC TTAAAGCTTA	1024
TTTTTATAAA GCTTTTTCAT AAAAAAAAAA AAAAAAAAAA AAAAAAAAAA AAA	1077

## (2) INFORMATION FOR SEQ ID NO:4:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 300 amino acids
- (B) TYPE: amino acid
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: protein

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:4:

Met Arg Ala Leu Glu Gly Pro Gly Leu Ser Leu Leu Cys Leu Val Leu  
 1 5 10 15  
 Ala Leu Pro Ala Leu Leu Pro Val Pro Ala Val Arg Gly Val Ala Glu  
 20 25 30  
 Thr Pro Thr Tyr Pro Trp Arg Asp Ala Glu Thr Gly Glu Arg Leu Val  
 35 40 45  
 Cys Ala Gln Cys Pro Pro Gly Thr Phe Val Gln Arg Pro Cys Arg Arg  
 50 55 60  
 Asp Ser Pro Thr Thr Cys Gly Pro Cys Pro Pro Arg His Tyr Thr Gln  
 65 70 75 80  
 Phe Trp Asn Tyr Leu Glu Arg Cys Arg Tyr Cys Asn Val Leu Cys Gly  
 85 90 95  
 Glu Arg Glu Glu Glu Ala Arg Ala Cys His Ala Thr His Asn Arg Ala  
 100 105 110  
 Cys Arg Cys Arg Thr Gly Phe Phe Ala His Ala Gly Phe Cys Leu Glu  
 115 120 125  
 His Ala Ser Cys Pro Pro Gly Ala Gly Val Ile Ala Pro Gly Thr Pro  
 130 135 140  
 Ser Gln Asn Thr Gln Cys Gln Pro Cys Pro Pro Gly Thr Phe Ser Ala  
 145 150 155 160  
 Ser Ser Ser Ser Ser Glu Gln Cys Gln Pro His Arg Asn Cys Thr Ala  
 165 170 175  
 Leu Gly Leu Ala Leu Asn Val Pro Gly Ser Ser Ser His Asp Thr Leu  
 180 185 190  
 Cys Thr Ser Cys Thr Gly Phe Pro Leu Ser Thr Arg Val Pro Gly Ala  
 195 200 205  
 Glu Glu Cys Glu Arg Ala Val Ile Asp Phe Val Ala Phe Gln Asp Ile  
 210 215 220  
 Ser Ile Lys Arg Leu Gln Arg Leu Leu Gln Ala Leu Glu Ala Pro Glu  
 225 230 235 240  
 Gly Trp Gly Pro Thr Pro Arg Ala Gly Arg Ala Ala Leu Gln Leu Lys  
 245 250 255  
 Leu Arg Arg Arg Leu Thr Glu Leu Leu Gly Ala Gln Asp Gly Ala Leu  
 260 265 270  
 Leu Val Arg Leu Leu Gln Ala Leu Arg Val Ala Arg Met Pro Gly Leu  
 275 280 285

Glu Arg Ser Val Arg Glu Arg Phe Leu Pro Val His  
 290 295 300

## (2) INFORMATION FOR SEQ ID NO:5:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 1667 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: DNA (genomic)

## (ix) FEATURE:

- (A) NAME/KEY: CDS
- (B) LOCATION: 73..582

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:5:

TGGCATGTCG GTCAGGCACA GCAGGGTCCT GTGTCCGCGC TGAGCCGCGC TCTCCCTGCT	60
CCAGCAAGGA CC ATG AGG GCG CTG GAG GGG CCA GGC CTG TCG CTG CTG	108
Met Arg Ala Leu Glu Gly Pro Gly Leu Ser Leu Leu	
1 5 10	
TGC CTG GTG TTG GCG CTG CCT GCC CTG CTG CCG GTG CCG GCT GTA CGC	156
Cys Leu Val Leu Ala Leu Pro Ala Leu Leu Pro Val Pro Ala Val Arg	
15 20 25	
GGA GTG GCA GAA ACA CCC ACC TAC CCC TGG CCG GAC GCA GAG ACA GGG	204
Gly Val Ala Glu Thr Pro Thr Tyr Pro Trp Arg Asp Ala Glu Thr Gly	
30 35 40	
GAG CCG CTG GTG TGC GCC CAG TGC CCC CCA GGC ACC TTT GTG CAG CCG	252
Glu Arg Leu Val Cys Ala Gln Cys Pro Pro Gly Thr Phe Val Gln Arg	
45 50 55 60	
CCG TGC CGC CGA GAC AGC CCC ACG ACG TGT GGC CCG TGT CCA CCG CGC	300
Pro Cys Arg Arg Asp Ser Pro Thr Thr Cys Gly Pro Cys Pro Pro Arg	
65 70 75	
CAC TAC ACG CAG TTC TGG AAC TAC CTG GAG CGC TGC CGC TAC TGC AAC	348
His Tyr Thr Gln Phe Trp Asn Tyr Leu Glu Arg Cys Arg Tyr Cys Asn	
80 85 90	
GTC CTC TGC GGG GAG CGT GAG GAG GAG GCA CCG GCT TGC CAC GCC ACC	396
Val Leu Cys Gly Glu Arg Glu Glu Glu Ala Arg Ala Cys His Ala Thr	
95 100 105	
CAC AAC CGT GCC TGC CGC TGC CGC ACC GGC TTC TTC GCG CAC GCT GGT	444
His Asn Arg Ala Cys Arg Cys Arg Thr Gly Phe Phe Ala His Ala Gly	
110 115 120	



TTC TGC TTG GAG CAC GCA TCG TGT CCA CCT GGT GCC GGC GTG ATT GCC	492
Phe Cys Leu Glu His Ala Ser Cys Pro Pro Gly Ala Gly Val Ile Ala	
125 130 135 140	
CCG GGT GAG AGC TGG GCG AGG GGA GGG GCC CCC AGG AGT GGT GGC CGG	540
Pro Gly Glu Ser Trp Ala Arg Gly Gly Ala Pro Arg Ser Gly Gly Arg	
145 150 155	
AGG TGT GGC AGG GGT CAG GTT GCT GGT CCC AGC CTT GCA CCC	582
Arg Cys Gly Arg Gly Gln Val Ala Gly Pro Ser Leu Ala Pro	
160 165 170	
TGAGCTAGGA CACCAGTTCC CTTGACCCTG TTCTTCCCTC CTGGCTGCAG GCACCCCCAG	642
CCAGAACACG CAGTGCCAGC CGTGCCCCC AGGCACCTTC TCAGCCAGCA GCTCCAGCTC	702
AGAGCAGTGC CAGGCCACC GCAACTGCAC GGCCCTGGGC CTGGCCCTCA ATGTGCCAGG	762
CTCTTCTCC CATGACACCC TGTGCACCAG CTGCACTGGC TTCCCTCTCA GCACCAGGT	822
ACCAGGTGAG CCAGAGGCCT GAGGGGGCAG CACACTGCAG GCCAGGCCCA CTTGTGCCCT	882
CACTCCTGCC CTTGCACGTG CATCTAGCCT GAGGCATGCC AGCTGGCTCT GGGAAGGGGC	942
CACAGTGGAT TTGAGGGGTC AGGGGTCCCT CCACTAGATC CCCACCAAGT CTGCCCTCTC	1002
AGGGGTGGCT GAGAATTGG ATCTGAGCCA GGGCAGGCC TCCCCTGGAG AGCTCTGGGA	1062
AAGTGGGCAG CAATCTCTTA ACTGCCCCGAG GGGAAGGTGG CTGGCTCCTC TGACACGGGG	1122
AAACCGAGGC CTGATGGTAA CTCTCCTAAC TGCCTGAGAG GAAGGTGGCT GCCTCCTCTG	1182
ACATGGGGAA ACCGAGGCC AATGTTAACC ACTGTTGAGA AGTCACAGGG GGAAGTGACC	1242
CCCTTAACAT CAAGTCAGGT CCGGTCCATC TGCAGGTCCC AACTCGCCCC TTCCGATGGC	1302
CCAGGAGCCC CAAGCCCTTG CTTGGGCCCC CTTGCCTCTT GCAGCCAAGG TCCGAGTGGC	1362
CGCTCCTGCC CCCTAGGCCT TTGCTCCAGC TCTCTGACCG AAGGCTCCTG CCCCTTCTCC	1422
AGTCCCCATC GTTGCACTGC CCTCTCCAGC ACGGCTCACT GCACAGGGAT TTCTCTCTCC	1482
TGCAAACCCC CCGAGTGGGG CCCAGAAAGC AGGGTACCTG GCAGCCCCCG CCAGTGTGTG	1542
TGGGTGAAAT GATCGGACCG CTGCCTCCCC ACCCACTGC AGGAGCTGAG GAGTGTGAGC	1602
GTGCCGTCAT CGACTTTGTG GCTTTCCAGG ACATCTCCAT CAAGAGGAGC GGCTGCTGCA	1662
GGCCC	1667

(2) INFORMATION FOR SEQ ID NO:6:

- (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 170 amino acids

(B) TYPE: amino acid  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:6:

```

Met Arg Ala Leu Glu Gly Pro Gly Leu Ser Leu Leu Cys Leu Val Leu
 1             5             10             15

Ala Leu Pro Ala Leu Leu Pro Val Pro Ala Val Arg Gly Val Ala Glu
 20             25             30

Thr Pro Thr Tyr Pro Trp Arg Asp Ala Glu Thr Gly Glu Arg Leu Val
 35             40             45

Cys Ala Gln Cys Pro Pro Gly Thr Phe Val Gln Arg Pro Cys Arg Arg
 50             55             60

Asp Ser Pro Thr Thr Cys Gly Pro Cys Pro Pro Arg His Tyr Thr Gln
 65             70             75             80

Phe Trp Asn Tyr Leu Glu Arg Cys Arg Tyr Cys Asn Val Leu Cys Gly
 85             90             95

Glu Arg Glu Glu Glu Ala Arg Ala Cys His Ala Thr His Asn Arg Ala
 100            105            110

Cys Arg Cys Arg Thr Gly Phe Phe Ala His Ala Gly Phe Cys Leu Glu
 115            120            125

His Ala Ser Cys Pro Pro Gly Ala Gly Val Ile Ala Pro Gly Glu Ser
 130            135            140

Trp Ala Arg Gly Gly Ala Pro Arg Ser Gly Gly Arg Arg Cys Gly Arg
 145            150            155            160

Gly Gln Val Ala Gly Pro Ser Leu Ala Pro
 165            170

```

(2) INFORMATION FOR SEQ ID NO:7:

(i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 455 amino acids  
 (B) TYPE: amino acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:7:

Met Gly Leu Ser Thr Val Pro Asp Leu Leu Leu Pro Leu Val Leu Leu  
 1 5 10 15  
 Glu Leu Leu Val Gly Ile Tyr Pro Ser Gly Val Ile Gly Leu Val Pro  
 20 25 30  
 His Leu Gly Asp Arg Glu Lys Arg Asp Ser Val Cys Pro Gln Gly Lys  
 35 40 45  
 Tyr Ile His Pro Gln Asn Asn Ser Ile Cys Cys Thr Lys Cys His Lys  
 50 55 60  
 Gly Thr Tyr Leu Tyr Asn Asp Cys Pro Gly Pro Gly Gln Asp Thr Asp  
 65 70 75 80  
 Cys Arg Glu Cys Glu Ser Gly Ser Phe Thr Ala Ser Glu Asn His Leu  
 85 90 95  
 Arg His Cys Leu Ser Cys Ser Lys Cys Arg Lys Glu Met Gly Gln Val  
 100 105 110  
 Glu Ile Ser Ser Cys Thr Val Asp Arg Asp Thr Val Cys Gly Cys Arg  
 115 120 125  
 Lys Asn Gln Tyr Arg His Tyr Trp Ser Glu Asn Leu Phe Gln Cys Phe  
 130 135 140  
 Asn Cys Ser Leu Cys Leu Asn Gly Thr Val His Leu Ser Cys Gln Glu  
 145 150 155 160  
 Lys Gln Asn Thr Val Cys Thr Cys His Ala Gly Phe Phe Leu Arg Glu  
 165 170 175  
 Asn Glu Cys Val Ser Cys Ser Asn Cys Lys Lys Ser Leu Glu Cys Thr  
 180 185 190  
 Lys Leu Cys Leu Pro Gln Ile Glu Asn Val Lys Gly Thr Glu Asp Ser  
 195 200 205  
 Gly Thr Thr Val Leu Leu Pro Leu Val Ile Phe Phe Gly Leu Cys Leu  
 210 215 220  
 Leu Ser Leu Leu Phe Ile Gly Leu Met Tyr Arg Tyr Gln Arg Trp Lys  
 225 230 235 240  
 Ser Lys Leu Tyr Ser Ile Val Cys Gly Lys Ser Thr Pro Glu Lys Glu  
 245 250 255  
 Gly Glu Leu Glu Gly Thr Thr Thr Lys Pro Leu Ala Pro Asn Pro Ser  
 260 265 270  
 Phe Ser Pro Thr Pro Gly Phe Thr Pro Thr Leu Gly Phe Ser Pro Val  
 275 280 285

Pro Ser Ser Thr Phe Thr Ser Ser Ser Thr Tyr Thr Pro Gly Asp Cys  
290 295 300

Pro Asn Phe Ala Ala Pro Arg Arg Glu Val Ala Pro Pro Tyr Gln Gly  
305 310 315 320

Ala Asp Pro Ile Leu Ala Thr Ala Leu Ala Ser Asp Pro Ile Pro Asn  
325 330 335

Pro Leu Gln Lys Trp Glu Asp Ser Ala His Lys Pro Gln Ser Leu Asp  
340 345 350

Thr Asp Asp Pro Ala Thr Leu Tyr Ala Val Val Glu Asn Val Pro Pro  
355 360 365

Leu Arg Trp Lys Glu Phe Val Arg Arg Leu Gly Leu Ser Asp His Glu  
370 375 380

Ile Asp Arg Leu Glu Leu Gln Asn Gly Arg Cys Leu Arg Glu Ala Gln  
385 390 395 400

Tyr Ser Met Leu Ala Thr Trp Arg Arg Arg Thr Pro Arg Arg Glu Ala  
405 410 415

Thr Leu Glu Leu Leu Gly Arg Val Leu Arg Asp Met Asp Leu Leu Gly  
420 425 430

Cys Leu Glu Asp Ile Glu Glu Ala Leu Cys Gly Pro Ala Ala Leu Pro  
435 440 445

Pro Ala Pro Ser Leu Leu Arg  
450 455

(2) INFORMATION FOR SEQ ID NO:8:

- (i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 461 amino acids  
(B) TYPE: amino acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:8:

Met Ala Pro Val Ala Val Trp Ala Ala Leu Ala Val Gly Leu Glu Leu  
1 5 10 15  
Trp Ala Ala Ala His Ala Leu Pro Ala Gln Val Ala Phe Thr Pro Tyr  
20 25 30

Ala Pro Glu Pro Gly Ser Thr Cys Arg Leu Arg Glu Tyr Tyr Asp Gln  
35 40 45

Thr Ala Gln Met Cys Cys Ser Lys Cys Ser Pro Gly Gln His Ala Lys  
50 55 60

Val Phe Cys Thr Lys Thr Ser Asp Thr Val Cys Asp Ser Cys Glu Asp  
65 70 75 80

Ser Thr Tyr Thr Gln Leu Trp Asn Trp Val Pro Glu Cys Leu Ser Cys  
85 90 95

Gly Ser Arg Cys Ser Ser Asp Gln Val Glu Thr Gln Ala Cys Thr ; 9  
100 105 110

Glu Gln Asn Arg Ile Cys Thr Cys Arg Pro Gly Trp Tyr Cys Ala Leu  
115 120 125

Ser Lys Gln Glu Gly Cys Arg Leu Cys Ala Pro Leu Arg Lys Cys Arg  
130 135 140

Pro Gly Phe Gly Val Ala Arg Pro Gly Thr Glu Thr Ser Asp Val Val  
145 150 155 160

Cys Lys Pro Cys Ala Pro Gly Thr Phe Ser Asn Thr Thr Ser Ser Thr  
165 170 175

Asp Ile Cys Arg Pro His Gln Ile Cys Asn Val Val Ala Ile Pro Gly  
180 185 190

Asn Ala Ser Arg Asp Ala Val Cys Thr Ser Thr Ser Pro Thr Arg Ser  
195 200 205

Met Ala Pro Gly Ala Val His Leu Pro Gln Pro Val Ser Thr Arg Ser  
210 215 220

Gln His Thr Gln Pro Thr Pro Glu Pro Ser Thr Ala Pro Ser Thr Ser  
225 230 235 240

Phe Leu Leu Pro Met Gly Pro Ser Pro Pro Ala Glu Gly Ser Thr Gly  
245 250 255

Asp Phe Ala Leu Pro Val Gly Leu Ile Val Gly Val Thr Ala Leu Gly  
260 265 270

Leu Leu Ile Ile Gly Val Val Asn Cys Val Ile Met Thr Gln Val Lys  
275 280 285

Lys Lys Pro Leu Cys Leu Gln Arg Glu Ala Lys Val Pro His Leu Pro  
290 295 300

Ala Asp Lys Ala Arg Gly Thr Gln Gly Pro Glu Gln Gln His Leu Leu  
305 310 315 320

Ile Thr Ala Pro Ser Ser Ser Ser Ser Leu Glu Ser Ser Ala Ser  
 325 330 335

Ala Leu Asp Arg Arg Ala Pro Thr Arg Asn Gln Pro Gln Ala Pro Gly  
 340 345 350

Val Glu Ala Ser Gly Ala Gly Glu Ala Arg Ala Ser Thr Gly Ser Ser  
 355 360 365

Asp Ser Ser Pro Gly Gly His Gly Thr Gln Val Asn Val Thr Cys Ile  
 370 375 380

Val Asn Val Cys Ser Ser Ser Asp His Ser Ser Gln Cys Ser Ser Gln  
 385 390 395 400

Ala Ser Ser Thr Met Gly Asp Thr Asp Ser Ser Pro Ser Glu Ser Pro  
 405 410 415

Lys Asp Glu Gln Val Pro Phe Ser Lys Glu Glu Cys Ala Phe Arg Ser  
 420 425 430

Gln Leu Glu Thr Pro Glu Thr Leu Leu Gly Ser Thr Glu Glu Ly Pro  
 435 440 445

Leu Pro Leu Gly Val Pro Asp Ala Gly Met Lys Pro Ser  
 450 455 460

## (2) INFORMATION FOR SEQ ID NO:9:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 427 amino acids
- (B) TYPE: amino acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: protein

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:9:

Met Gly Ala Gly Ala Thr Gly Arg Ala Met Asp Gly Pro Arg Leu Leu  
 1 5 10 15

Leu Leu Leu Leu Leu Gly Val Ser Leu Gly Gly Ala Lys Glu Ala Cys  
 20 25 30

Pro Thr Gly Leu Tyr Thr His Ser Gly Glu Cys Cys Lys Ala Cys Asn  
 35 40 45

Leu Gly Glu Gly Val Ala Gln Pro Cys Gly Ala Asn Gln Thr Val Cys  
 50 55 60

Glu Pro Cys Leu Asp Ser Val Thr Phe Ser Asp Val Val Ser Ala Thr  
 65 70 75 80  
 Glu Pro Cys Lys Pro Cys Thr Glu Cys Val Gly Leu Gln Ser Met Ser  
 85 90 95  
 Ala Pro Cys Val Glu Ala Asp Asp Ala Val Cys Arg Cys Ala Tyr Gly  
 100 105 110  
 Tyr Tyr Gln Asp Glu Thr Thr Gly Arg Cys Glu Ala Cys Arg Val Cys  
 115 120 125  
 Glu Ala Gly Ser Gly Leu Val Phe Ser Cys Gln Asp Lys Gln Asn Thr  
 130 135 140  
 Val Cys Glu Glu Cys Pro Asp Gly Thr Tyr Ser Asp Glu Ala Asn His  
 145 150 155 160  
 Val Asp Pro Cys Leu Pro Cys Thr Val Cys Glu Asp Thr Glu Arg Gln  
 165 170 175  
 Leu Arg Glu Cys Thr Arg Trp Ala Asp Ala Glu Cys Glu Glu Ile Pro  
 180 185 190  
 Gly Arg Trp Ile Thr Arg Ser Thr Pro Pro Glu Gly Ser Asp Ser Thr  
 195 200 205  
 Ala Pro Ser Thr Gln Glu Pro Glu Ala Pro Pro Glu Gln Asp Leu Ile  
 210 215 220  
 Ala Ser Thr Val Ala Gly Val Val Thr Thr Val Met Gly Ser Ser Gln  
 225 230 235 240  
 Pro Val Val Thr Arg Gly Thr Thr Asp Asn Leu Ile Pro Val Tyr Cys  
 245 250 255  
 Ser Ile Leu Ala Ala Val Val Val Gly Leu Val Ala Tyr Ile Ala Phe  
 260 265 270  
 Lys Arg Trp Asn Ser Cys Lys Gln Asn Lys Gln Gly Ala Asn Ser Arg  
 275 280 285  
 Pro Val Asn Gln Thr Pro Pro Pro Glu Gly Glu Lys Leu His Ser Asp  
 290 295 300  
 Ser Gly Ile Ser Val Asp Ser Gln Ser Leu His Asp Gln Gln Pro His  
 305 310 315 320  
 Thr Gln Thr Ala Ser Gly Gln Ala Leu Lys Gly Asp Gly Gly Leu Tyr  
 325 330 335  
 Ser Ser Leu Pro Pro Ala Lys Arg Glu Glu Val Glu Lys Leu Leu Asn  
 340 345 350

Gly Ser Ala Gly Asp Thr Trp Arg His Leu Ala Gly Glu Leu Gly Tyr  
 355 360 365

Gln Pro Glu His Ile Asp Ser Phe Thr His Glu Ala Cys Pro Val Arg  
 370 375 380

Ala Leu Leu Ala Ser Trp Ala Thr Gln Asp Ser Ala Thr Leu Asp Ala  
 385 390 395 400

Leu Leu Ala Ala Leu Arg Arg Ile Gln Arg Ala Asp Leu Val Glu Ser  
 405 410 415

Leu Cys Ser Glu Ser Thr Ala Thr Ser Pro Val  
 420 425

## (2) INFORMATION FOR SEQ ID NO:10:

- (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 415 amino acids  
 (B) TYPE: amino acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:10:

Met Arg Leu Pro Arg Ala Ser Ser Pro Cys Gly Leu Ala Trp Gly Pro  
 1 5 10 15

Leu Leu Leu Gly Leu Ser Gly Leu Leu Val Ala Ser Gln Pro Gln Leu  
 20 25 30

Val Pro Pro Tyr Arg Ile Glu Asn Gln Thr Cys Trp Asp Gln Asp Lys  
 35 40 45

Glu Tyr Tyr Glu Pro Met His Asp Val Cys Cys Ser Arg Cys Pro Pro  
 50 55 60

Gly Glu Phe Val Phe Ala Val Cys Ser Arg Ser Gln Asp Thr Val Cys  
 65 70 75 80

Lys Thr Cys Pro His Asn Ser Tyr Asn Glu His Trp Asn His Leu Ser  
 85 90 95

Thr Cys Gln Leu Cys Arg Pro Cys Asp Ile Val Leu Gly Phe Glu Glu  
 100 105 110

Val Ala Pro Cys Thr Ser Asp Arg Lys Ala Glu Cys Arg Cys Gln Pro  
 115 120 125



Gly Met Ser Cys Val Tyr Leu Asp Asn Glu Cys Val His Cys Glu Glu  
130 135 140

Glu Arg Leu Val Leu Cys Gln Pro Gly Thr Glu Ala Glu Val Thr Asp  
145 150 155 160

Glu Ile Met Asp Thr Asp Val Asn Cys Val Pro Cys Lys Pro Gly His  
165 170 175

Phe Gln Asn Thr Ser Ser Pro Arg Ala Arg Cys Gln Pro His Thr Arg  
180 185 190

Cys Glu Ile Gln Gly Leu Val Glu Ala Ala Pro Gly Thr Ser Tyr Ser  
195 200 205

Asp Thr Ile Cys Lys Asn Pro Pro Glu Pro Gly Ala Met Leu Leu Leu  
210 215 220

Ala Ile Leu Leu Ser Leu Val Leu Phe Leu Leu Phe Thr Thr Val Leu  
225 230 235 240

Ala Cys Ala Trp Met Arg His Pro Ser Leu Cys Arg Lys Leu Gly Thr  
245 250 255

Leu Leu Lys Arg His Pro Glu Gly Glu Glu Ser Pro Pro Cys Pro Ala  
260 265 270

Pro Arg Ala Asp Pro His Phe Pro Asp Leu Ala Glu Pro Leu Leu Pro  
275 280 285

Met Ser Gly Asp Leu Ser Pro Ser Pro Ala Gly Pro Pro Thr Ala Pro  
290 295 300

Ser Leu Glu Glu Val Val Leu Gln Gln Gln Ser Pro Leu Val Gln Ala  
305 310 315 320

Arg Glu Leu Glu Ala Glu Pro Gly Glu His Gly Gln Val Ala His Gly  
325 330 335

Ala Asn Gly Ile His Val Thr Gly Gly Ser Val Thr Val Thr Gly Asn  
340 345 350

Ile Tyr Ile Tyr Asn Gly Pro Val Leu Gly Gly Thr Arg Gly Pro Gly  
355 360 365

Asp Pro Pro Ala Pro Pro Glu Pro Pro Tyr Pro Thr Pro Glu Glu Gly  
370 375 380

Ala Pro Gly Pro Ser Glu Leu Ser Thr Pro Tyr Gln Glu Asp Gly Lys  
385 390 395 400

Ala Trp His Leu Ala Glu Thr Glu Thr Leu Gly Cys Gln Asp Leu  
405 410 415

(2) INFORMATION FOR SEQ ID NO:11:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 335 amino acids
- (B) TYPE: amino acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:11:

```

Met Leu Gly Ile Trp Thr Leu Leu Pro Leu Val Leu Thr Ser Val Ala
1           5           10           15

Arg Leu Ser Ser Lys Ser Val Asn Ala Gln Val Thr Asp Ile Asn Ser
          20           25           30

Lys Gly Leu Glu Leu Arg Lys Thr Val Thr Thr Val Glu Thr Gln Asn
          35           40           45

Leu Glu Gly Leu His His Asp Gly Gln Phe Cys His Lys Pro Cys Pro
          50           55           60

Pro Gly Glu Arg Lys Ala Arg Asp Cys Thr Val Asn Gly Asp Glu Pro
          65           70           75           80

Asp Cys Val Pro Cys Gln Glu Gly Lys Glu Tyr Thr Asp Lys Ala His
          85           90           95

Phe Ser Ser Lys Cys Arg Arg Cys Arg Leu Cys Asp Glu Gly His Gly
          100          105          110

Leu Glu Val Glu Ile Asn Cys Thr Arg Thr Gln Asn Thr Lys Cys Arg
          115          120          125

Cys Lys Pro Asn Phe Phe Cys Asn Ser Thr Val Cys Glu His Cys Asp
          130          135          140

Pro Cys Thr Lys Cys Glu His Gly Ile Ile Lys Glu Cys Thr Leu Thr
          145          150          155          160

Ser Asn Thr Lys Cys Lys Glu Glu Gly Ser Arg Ser Asn Leu Gly Trp
          165          170          175

Leu Cys Leu Leu Leu Leu Pro Ile Pro Leu Ile Val Trp Val Lys Arg
          180          185          190

Lys Glu Val Gln Lys Thr Cys Arg Lys His Arg Lys Glu Asn Gln Gly
          195          200          205

Ser His Glu Ser Pro Thr Leu Asn Pro Glu Thr Val Ala Ile Asn Leu
          210          215          220

```

Ser Asp Val Asp Leu Ser Lys Tyr Ile Thr Thr Ile Ala Gly Val Met  
 225 230 235 240  
 Thr Leu Ser Gln Val Lys Gly Phe Val Arg Lys Asn Gly Val Asn Glu  
 245 250 255  
 Ala Lys Ile Asp Glu Ile Lys Asn Asp Asn Val Gln Asp Thr Ala Glu  
 260 265 270  
 Gln Lys Val Gln Leu Leu Arg Asn Trp His Gln Leu His Gly Lys Lys  
 275 280 285  
 Glu Ala Tyr Asp Thr Leu Ile Lys Asp Leu Lys Lys Ala Asn Leu Cys  
 290 295 300  
 Thr Leu Ala Glu Lys Ile Gln Thr Ile Ile Leu Lys Asp Ile Thr Ser  
 305 310 315 320  
 Asp Ser Glu Asn Ser Asn Phe Arg Asn Glu Ile Gln Ser Leu Val  
 325 330 335

## (2) INFORMATION FOR SEQ ID NO:12:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 260 amino acids
  - (B) TYPE: amino acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:12:

Met Ala Arg Pro His Pro Trp Trp Leu Cys Val Leu Gly Thr Leu Val  
 1 5 10 15  
 Gly Leu Ser Ala Thr Pro Ala Pro Lys Ser Cys Pro Glu Arg His Tyr  
 20 25 30  
 Trp Ala Gln Gly Lys Leu Cys Cys Gln Met Cys Glu Pro Gly Thr Phe  
 35 40 45  
 Leu Val Lys Asp Cys Asp Gln His Arg Lys Ala Ala Gln Cys Asp Pro  
 50 55 60  
 Cys Ile Pro Gly Val Ser Phe Ser Pro Asp His His Thr Arg Pro His  
 65 70 75 80  
 Cys Glu Ser Cys Arg His Cys Asn Ser Gly Leu Leu Val Arg Asn Cys  
 85 90 95

Thr Ile Thr Ala Asn Ala Glu Cys Ala Cys Arg Asn Gly Trp Gln Cys  
 100 105 110  
 Arg Asp Lys Glu Cys Thr Glu Cys Asp Pro Leu Pro Asn Pro Ser Leu  
 115 120 125  
 Thr Ala Arg Ser Ser Gln Ala Leu Ser Pro His Pro Gln Pro Thr His  
 130 135 140  
 Leu Pro Tyr Val Ser Glu Met Leu Glu Ala Arg Thr Ala Gly His Met  
 145 150 155 160  
 Gln Thr Leu Ala Asp Phe Arg Gln Leu Pro Ala Arg Thr Leu Ser Thr  
 165 170 175  
 His Trp Pro Pro Gln Arg Ser Leu Cys Ser Ser Asp Phe Ile Arg Ile  
 180 185 190  
 Leu Val Ile Phe Ser Gly Met Phe Leu Val Phe Thr Leu Ala Gly Ala  
 195 200 205  
 Leu Phe Leu His Gln Arg Arg Lys Tyr Arg Ser Asn Lys Gly Glu Ser  
 210 215 220  
 Pro Val Glu Pro Ala Glu Pro Cys Arg Tyr Ser Cys Pro Arg Glu Glu  
 225 230 235 240  
 Glu Gly Ser Thr Ile Pro Ile Gln Glu Asp Tyr Arg Lys Pro Glu Pro  
 245 250 255  
 Ala Cys Ser Pro  
 260

## (2) INFORMATION FOR SEQ ID NO:13:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 595 amino acids
  - (B) TYPE: amino acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:13:

Met Arg Val Leu Leu Ala Ala Leu Gly Leu Leu Phe Leu Gly Ala Leu  
 1 5 10 15  
 Arg Ala Phe Pro Gln Asp Arg Pro Phe Glu Asp Thr Cys His Gly Asn  
 20 25 30

Pro Ser His Tyr Tyr Asp Lys Ala Val Arg Arg Cys Cys Tyr Arg Cys  
 35 40 45  
 Pro Met Gly Leu Phe Pro Thr Gln Gln Cys Pro Gln Arg Pro Thr Asp  
 50 55 60  
 Cys Arg Lys Gln Cys Glu Pro Asp Tyr Tyr Leu Asp Glu Ala Asp Arg  
 65 70 75 80  
 Cys Thr Ala Cys Val Thr Cys Ser Arg Asp Asp Leu Val Glu Lys Thr  
 85 90 95  
 Pro Cys Ala Trp Asn Ser Ser Arg Val Cys Glu Cys Arg Pro Gly Met  
 100 105 110  
 Phe Cys Ser Thr Ser Ala Val Asn Ser Cys Ala Arg Cys Phe Phe His  
 115 120 125  
 Ser Val Cys Pro Ala Gly Met Ile Val Lys Phe Pro Gly Thr Ala Gln  
 130 135 140  
 Lys Asn Thr Val Cys Glu Pro Ala Ser Pro Gly Val Ser Pro Ala Cys  
 145 150 155 160  
 Ala Ser Pro Glu Asn Cys Lys Glu Pro Ser Ser Gly Thr Ile Pro Gln  
 165 170 175  
 Ala Lys Pro Thr Pro Val Ser Pro Ala Thr Ser Ser Ala Ser Thr Met  
 180 185 190  
 Pro Val Arg Gly Gly Thr Arg Leu Ala Gln Glu Ala Ala Ser Lys Leu  
 195 200 205  
 Thr Arg Ala Pro Asp Ser Pro Ser Ser Val Gly Arg Pro Ser Ser Asp  
 210 215 220  
 Pro Gly Leu Ser Pro Thr Gln Pro Cys Pro Glu Gly Ser Gly Asp Cys  
 225 230 235 240  
 Arg Lys Gln Cys Glu Pro Asp Tyr Tyr Leu Asp Glu Ala Gly Arg Cys  
 245 250 255  
 Thr Ala Cys Val Ser Cys Ser Arg Asp Asp Leu Val Glu Lys Thr Pro  
 260 265 270  
 Cys Ala Trp Asn Ser Ser Arg Thr Cys Glu Cys Arg Pro Gly Met Ile  
 275 280 285  
 Cys Ala Thr Ser Ala Thr Asn Ser Cys Ala Arg Cys Val Pro Tyr Pro  
 290 295 300  
 Ile Cys Ala Ala Glu Thr Val Thr Lys Pro Gln Asp Met Ala Glu Lys  
 305 310 315 320

Asp Thr Thr Phe Glu Ala Pro Pro Leu Gly Thr Gln Pro Asp Cys Asn  
 325 330 335  
 Pro Thr Pro Glu Asn Gly Glu Ala Pro Ala Ser Thr Ser Pro Thr Gln  
 340 345 350  
 Ser Leu Leu Val Asp Ser Gln Ala Ser Lys Thr Leu Pro Ile Pro Thr  
 355 360 365  
 Ser Ala Pro Val Ala Leu Ser Ser Thr Gly Lys Pro Val Leu Asp Ala  
 370 375 380  
 Gly Pro Val Leu Phe Trp Val Ile Leu Val Leu Val Val Val Gly  
 385 390 395 400  
 Ser Ser Ala Phe Leu Leu Cys His Arg Arg Ala Cys Arg Lys Arg Ile  
 405 410 415  
 Arg Gln Lys Leu His Leu Cys Tyr Pro Val Gln Thr Ser Gln Pro Lys  
 420 425 430  
 Leu Glu Leu Val Asp Ser Arg Pro Arg Arg Ser Ser Thr Gln Leu Arg  
 435 440 445  
 Ser Gly Ala Ser Val Thr Glu Pro Val Ala Glu Glu Arg Gly Leu Met  
 450 455 460  
 Ser Gln Pro Leu Met Glu Thr Cys His Ser Val Gly Ala Ala Tyr Leu  
 465 470 475 480  
 Glu Ser Leu Pro Leu Gln Asp Ala Ser Pro Ala Gly Gly Pro Ser Ser  
 485 490 495  
 Pro Arg Asp Leu Pro Glu Pro Arg Val Ser Thr Glu His Thr Asn Asn  
 500 505 510  
 Lys Ile Glu Lys Ile Tyr Ile Met Lys Ala Asp Thr Val Ile Val Gly  
 515 520 525  
 Thr Val Lys Ala Glu Leu Pro Glu Gly Arg Gly Leu Ala Gly Pro Ala  
 530 535 540  
 Glu Pro Glu Leu Glu Glu Glu Leu Glu Ala Asp His Thr Pro His Tyr  
 545 550 555 560  
 Pro Glu Gln Glu Thr Glu Pro Pro Leu Gly Ser Cys Ser Asp Val Met  
 565 570 575  
 Leu Ser Val Glu Glu Glu Gly Lys Glu Asp Pro Leu Pro Thr Ala Ala  
 580 585 590  
 Ser Gly Lys  
 595

(2) INFORMATION FOR SEQ ID NO:14:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 277 amino acids
- (B) TYPE: amino acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:14:

```

Met Val Arg Leu Pro Leu Gln Cys Val Leu Trp Gly Cys Leu Leu Thr
1           5           10           15

Ala Val His Pro Glu Pro Pro Thr Ala Cys Arg Glu Lys Gln Tyr Leu
20           25           30

Ile Asn Ser Gln Cys Cys Ser Leu Cys Gln Pro Gly Gln Lys Leu Val
35           40           45

Ser Asp Cys Thr Glu Phe Thr Glu Thr Glu Cys Leu Pro Cys Gly Glu
50           55           60

Ser Glu Phe Leu Asp Thr Trp Asn Arg Glu Thr His Cys His Gln His
65           70           75           80

Lys Tyr Cys Asp Pro Asn Leu Gly Leu Arg Val Gln Gln Lys Gly Thr
85           90           95

Ser Glu Thr Asp Thr Ile Cys Thr Cys Glu Glu Gly Trp His Cys Thr
100          105          110

Ser Glu Ala Cys Glu Ser Cys Val Leu His Arg Ser Cys Ser Pro Gly
115          120          125

Phe Gly Val Lys Gln Ile Ala Thr Gly Val Ser Asp Thr Ile Cys Glu
130          135          140

Pro Cys Pro Val Gly Phe Phe Ser Asn Val Ser Ser Ala Phe Glu Lys
145          150          155          160

Cys His Pro Trp Thr Ser Cys Glu Thr Lys Asp Leu Val Val Gln Gln
165          170          175

Ala Gly Thr Asn Lys Thr Asp Val Val Cys Gly Pro Gln Asp Arg Leu
180          185          190

Arg Ala Leu Val Val Ile Pro Ile Ile Phe Gly Ile Leu Phe Ala Ile
195          200          205

Leu Leu Val Leu Val Phe Ile Lys Lys Val Ala Lys Lys Pro Thr Asn
210          215          220

```

Lys Ala Pro His Pro Lys Gln Glu Pro Gln Glu Ile Asn Phe Pro Asp  
 225 230 235 240  
 Asp Leu Pro Gly Ser Asn Thr Ala Ala Pro Val Gln Glu Thr Leu His  
 245 250 255  
 Gly Cys Gln Pro Val Thr Gln Glu Asp Gly Lys Glu Ser Arg Ile Ser  
 260 265 270  
 Val Gln Glu Arg Gln  
 275

## (2) INFORMATION FOR SEQ ID NO:15:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 255 amino acids
- (B) TYPE: amino acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: protein

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:15:

Met Gly Asn Ser Cys Tyr Asn Ile Val Ala Thr Leu Leu Leu Val Leu  
 1 5 10 15  
 Asn Phe Glu Arg Thr Arg Ser Leu Gln Asp Pro Cys Ser Asn Cys Pro  
 20 25 30  
 Ala Gly Thr Phe Cys Asp Asn Asn Arg Asn Gln Ile Cys Ser Pro Cys  
 35 40 45  
 Pro Pro Asn Ser Phe Ser Ser Ala Gly Gly Gln Arg Thr Cys Asp Ile  
 50 55 60  
 Cys Arg Gln Cys Lys Gly Val Phe Arg Thr Arg Lys Glu Cys Ser Ser  
 65 70 75 80  
 Thr Ser Asn Ala Glu Cys Asp Cys Thr Pro Gly Phe His Cys Leu Gly  
 85 90 95  
 Ala Gly Cys Ser Met Cys Glu Gln Asp Cys Lys Gln Gly Gln Glu Leu  
 100 105 110  
 Thr Lys Lys Gly Cys Lys Asp Cys Cys Phe Gly Thr Phe Asn Asp Gln  
 115 120 125  
 Lys Arg Gly Ile Cys Arg Pro Trp Thr Asn Cys Ser Leu Asp Gly Lys  
 130 135 140



Ser Val Leu Val Asn Gly Thr Lys Glu Arg Asp Val Val Cys Gly Pro  
 145 150 155 160

Ser Pro Ala Asp Leu Ser Pro Gly Ala Ser Ser Val Thr Pro Pro Ala  
 165 170 175

Pro Ala Arg Glu Pro Gly His Ser Pro Gln Ile Ile Ser Phe Phe Leu  
 180 185 190

Ala Leu Thr Ser Thr Ala Leu Leu Phe Leu Leu Phe Phe Leu Thr Leu  
 195 200 205

Arg Phe Ser Val Val Lys Arg Gly Arg Lys Lys Leu Leu Tyr Ile Phe  
 210 215 220

Lys Gln Pro Phe Met Arg Pro Val Gln Thr Thr Gln Glu Glu Asp Gly  
 225 230 235 240

Cys Ser Cys Arg Phe Pro Glu Glu Glu Glu Gly Gly Cys Glu Leu  
 245 250 255

## (2) INFORMATION FOR SEQ ID NO:16:

- (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 277 amino acids  
 (B) TYPE: amino acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:16:

Met Cys Val Gly Ala Arg Arg Leu Gly Arg Gly Pro Cys Ala Ala Leu  
 1 5 10 15

Leu Leu Leu Gly Leu Gly Leu Ser Thr Val Thr Gly Leu His Cys Val  
 20 25 30

Gly Asp Thr Tyr Pro Ser Asn Asp Arg Cys Cys His Glu Cys Arg Pro  
 35 40 45

Gly Asn Gly Met Val Ser Arg Cys Ser Arg Ser Gln Asn Thr Val Cys  
 50 55 60

Arg Pro Cys Gly Pro Gly Phe Tyr Asn Asp Val Val Ser Ser Lys Pro  
 65 70 75 80

Cys Lys Pro Cys Thr Trp Cys Asn Leu Arg Ser Gly Ser Glu Arg Lys  
 85 90 95

Gln Leu Cys Thr Ala Thr Gln Asp Thr Val Cys Arg Cys Arg Ala Gly  
 100 105 110  
 Thr Gln Pro Leu Asp Ser Tyr Lys Pro Gly Val Asp Cys Ala Pro Cys  
 115 120 125  
 Pro Pro Gly His Phe Ser Pro Gly Asp Asn Gln Ala Cys Lys Pro Trp  
 130 135 140  
 Thr Asn Cys Thr Leu Ala Gly Lys His Thr Leu Gln Pro Ala Ser Asn  
 145 150 155 160  
 Ser Ser Asp Ala Ile Cys Glu Asp Arg Asp Pro Pro Ala Thr Gln Pro  
 165 170 175  
 Gln Glu Thr Gln Gly Pro Pro Ala Arg Pro Ile Thr Val Gln Pro Thr  
 180 185 190  
 Glu Ala Trp Pro Arg Thr Ser Gln Gly Pro Ser Thr Arg Pro Val Glu  
 195 200 205  
 Val Pro Gly Gly Arg Ala Val Ala Ala Ile Leu Gly Leu Gly Leu Val  
 210 215 220  
 Leu Gly Leu Leu Gly Pro Leu Ala Ile Leu Leu Ala Leu Tyr Leu Leu  
 225 230 235 240  
 Arg Arg Asp Gln Arg Leu Pro Pro Asp Ala His Lys Pro Pro Gly Gly  
 245 250 255  
 Gly Ser Phe Arg Thr Pro Ile Gln Glu Glu Gln Ala Asp Ala His Ser  
 260 265 270  
 Thr Leu Ala Lys Ile  
 275

## (2) INFORMATION FOR SEQ ID NO:17:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 349 amino acids
  - (B) TYPE: amino acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:17:

Met Lys Ser Val Leu Tyr Leu Tyr Ile Leu Phe Leu Ser Cys Ile Ile  
 1 5 10 15

Ile Asn Gly Arg Asp Ala Ala Pro Tyr Thr Pro Pro Asn Gly Lys Cys  
 20 25 30  
 Lys Asp Thr Glu Tyr Lys Arg His Asn Leu Cys Cys Leu Ser Cys Pro  
 35 40 45  
 Pro Gly Thr Tyr Ala Ser Arg Leu Cys Asp Ser Lys Thr Asn Thr Gln  
 50 55 60  
 Cys Thr Pro Cys Gly Ser Gly Thr Phe Thr Ser Arg Asn Asn His Leu  
 65 70 75 80  
 Pro Ala Cys Leu Ser Cys Asn Gly Arg Cys Asn Ser Asn Gln Val Glu  
 85 90 95  
 Thr Arg Ser Cys Asn Thr Thr His Asn Arg Ile Cys Glu Cys Ser Pro  
 100 105 110  
 Gly Tyr Tyr Cys Leu Leu Lys Gly Ser Ser Gly Cys Lys Ala Cys Val  
 115 120 125  
 Ser Gln Thr Lys Cys Gly Ile Gly Tyr Gly Val Ser Gly His Thr Ser  
 130 135 140  
 Val Gly Asp Val Ile Cys Ser Pro Cys Gly Phe Gly Thr Tyr Ser His  
 145 150 155 160  
 Thr Val Ser Ser Ala Asp Lys Cys Glu Pro Val Pro Asn Asn Thr Phe  
 165 170 175  
 Asn Tyr Ile Asp Val Glu Ile Thr Leu Tyr Pro Val Asn Asp Thr Ser  
 180 185 190  
 Cys Thr Arg Thr Thr Thr Thr Gly Leu Ser Glu Ser Ile Leu Thr Ser  
 195 200 205  
 Glu Leu Thr Ile Thr Met Asn His Thr Asp Cys Asn Pro Val Phe Arg  
 210 215 220  
 Glu Glu Tyr Phe Ser Val Leu Asn Lys Val Ala Thr Ser Gly Phe Phe  
 225 230 235 240  
 Thr Gly Glu Asn Arg Tyr Gln Asn Ile Ser Lys Val Cys Thr Leu Asn  
 245 250 255  
 Phe Glu Ile Lys Cys Asn Asn Lys Gly Ser Ser Phe Lys Gln Leu Thr  
 260 265 270  
 Lys Ala Lys Asn Asp Asp Gly Met Met Ser His Ser Glu Thr Val Thr  
 275 280 285  
 Leu Ala Gly Asp Cys Leu Ser Ser Val Asp Ile Tyr Ile Leu Tyr Ser  
 290 295 300

Asn Thr Asn Ala Gln Asp Tyr Glu Thr Asp Thr Ile Ser Tyr Arg Val  
305 310 315 320

Gly Asn Val Leu Asp Asp Asp Ser His Met Pro Gly Ser Cys Asn Ile  
325 330 335

His Lys Pro Ile Thr Asn Ser Lys Pro Thr Arg Phe Leu  
340 345

(2) INFORMATION FOR SEQ ID NO:18:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 355 amino acids
- (B) TYPE: amino acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:18:

Met Lys Ser Tyr Ile Leu Leu Leu Leu Ser Cys Ile Ile Ile Ile  
1 5 10 15

Asn Ser Asp Ile Thr Pro His Glu Pro Ser Asn Gly Lys Cys Lys Asp  
20 25 30

Asn Glu Tyr Lys Arg His His Leu Cys Cys Leu Ser Cys Pro Pro Gly  
35 40 45

Thr Tyr Ala Ser Arg Leu Cys Asp Ser Lys Thr Asn Thr Asn Thr Gln  
50 55 60

Cys Thr Pro Cys Ala Ser Asp Thr Phe Thr Ser Arg Asn Asn His Leu  
65 70 75 80

Pro Ala Cys Leu Ser Cys Asn Gly Arg Cys Asp Ser Asn Gln Val Glu  
85 90 95

Thr Arg Ser Cys Asn Thr Thr His Asn Arg Ile Cys Asp Cys Ala Pro  
100 105 110

Gly Tyr Tyr Cys Phe Leu Lys Gly Ser Ser Gly Cys Lys Ala Cys Val  
115 120 125

Ser Gln Thr Lys Cys Gly Ile Gly Tyr Gly Val Ser Gly His Thr Pro  
130 135 140

Thr Gly Asp Val Val Cys Ser Pro Cys Gly Leu Gly Thr Thr Ser His  
145 150 155 160

Thr Val Ser Ser Val Asp Lys Cys Glu Pro Val Pro Ser Asn Thr Phe  
 165 170 175  
 Asn Tyr Ile Asp Val Glu Ile Asn Leu Tyr Pro Val Asn Asp Thr Ser  
 180 185 190  
 Cys Thr Arg Thr Thr Thr Thr Gly Leu Ser Glu Ser Ile Ser Thr Ser  
 195 200 205  
 Glu Leu Thr Ile Thr Met Asn His Lys Asp Cys Asp Pro Val Phe Arg  
 210 215 220  
 Asn Gly Tyr Phe Ser Val Leu Asn Glu Val Ala Thr Ser Gly Phe Phe  
 225 230 235 240  
 Thr Gly Gln Asn Arg Tyr Gln Asn Ile Ser Lys Val Cys Thr Leu Asn  
 245 250 255  
 Phe Glu Ile Lys Cys Asn Asn Lys Asp Ser Tyr Ser Ser Ser Lys Gln  
 260 265 270  
 Leu Thr Lys Thr Lys Asn Asp Asp Asp Ser Ile Met Pro His Ser Glu  
 275 280 285  
 Ser Val Thr Leu Val Gly Asp Cys Leu Ser Ser Val Asp Ile Tyr Ile  
 290 295 300  
 Leu Tyr Ser Asn Thr Asn Thr Gln Asp Tyr Glu Thr Asp Thr Ile Ser  
 305 310 315 320  
 Tyr His Val Gly Asn Val Leu Asp Val Asp Ser His Met Pro Gly Arg  
 325 330 335  
 Cys Asp Thr His Lys Leu Ile Thr Asn Ser Asn Ser Gln Tyr Pro Thr  
 340 345 350  
 His Phe Leu  
 355

## (2) INFORMATION FOR SEQ ID NO:19:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 506 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:19:

GAATTCGGCA NAGCCTCTCC ACGCGCAGAA CTCAGCCAAC GATTCTGAT AGATTTTGG

GAGTTTGACC AGAGATGCAA GGGGTGAAGG AGCGCTTCCT ACCGTTAGGA ACTCTGGGGA	120
CAGNNGCCCC CGGCCGCTG ATGGCCGAGG CAGGGTGCGA CCCAGGACCC AGGACGGCGT	180
CGGGAACCAT ACCATGGCCC GGATCCCCAA GACCCTAAAG TTCGTGGTCG TCATCGTCGC	240
GGTCCTGCTG CCAGTCCTAG CTTACTCTGC CACCACTGCC CGGCAGAGGA AGTTNCCAG	300
CAGNCANTCG NCCCACAGCA ACAGNGGCAC AGTTTCAAGG GGGNAGGAGT TTTCCANCAA	360
GTTTTTATAG TTCAGACNT ATTGGNGCTN TNAACCCTTG CACAAGGGTT TGGNTTAAAC	420
CAANGTTTCC AANATGNACT TTTTNGTTCC CTGTTANATT TTTTAATTAG TTNAANTTAA	480
ATTINTNAAC CTNCCNGGG NAAATT	506

## (2) INFORMATION FOR SEQ ID NO:20:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 325 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:20:

GGCAGAGGTG TCTCCAGCCT GGCTCTATCT TCCTCCTTGT NATCGTCCCA TCCCCACATC	60
CCGTGCAC : CCCAGGACCC TGGTCTCATC AGTCCCTCTC CTGGAGCTGG GGGTCCACAC	120
ATCTCCCAGC CAAGTCCAAG AGGGCAGGGC CAGTTCCTCC CATCTTCAGG CCCAGCCAGG	180
CAGGGGGCAG TCGGCTCCTC AACTGGGTGA CAAGGGTGAG GATGAGAAGT GGTCACGGGG	240
ATTTATTAG CTTGGTCAG AGCAGAACAC AGATTTTCC GTGTGTTGGT TTTTACTCTN	300
NTTCCCCTTC TTNATNCCCC TTTCN	325

## (2) INFORMATION FOR SEQ ID NO:21:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 340 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:21:

GGCAGAGGCC CCAGCTGCTG AAGAGACAAT AATCACCAGC CCGGGGACTC CTGNNTCTNC 60  
 TNATTACCTC TNATGCACCA TCGTAGGGAT CATAGTTCTA ATTGTGCCTT CTAATTGTTT 120  
 TTGTTTGAAA AGANTTCACT GTGGAAGAAA TTCCTTCCTT ACCTGTAAGT TNCAGGTAGG 180  
 NGCCTGGCTG AGGGCGGGGG GCGCTGGTAC ACTCTGTGAC CCTGCCTCCC TCTGNCTGTT 240  
 TTCCACAGA CAGAAACGCC TGCCCTGNC CCCAAGTTC TNGTGTTTT CAGCCTGGCT 300  
 CTATCTTNNC TCCTTGTA TCGTTCCCAT CCCCACANGC 340

## (2) INFORMATION FOR SEQ ID NO:22:

- (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 241 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:22:

CCAGGGTCTC CTNCCCCACC TGCTGAAGAG ACANTGACCA CCAGCCCGGG GACTCCTGCC 60  
 TCTTCCTCAT TACCTCTNAT GNANCATCGT AGGGATCATA GTTCTAATTG TGCCTTCTGA 120  
 ATTGTGCTTT GTTTGAAAG ACTTCACTGT GGAAGAAAT TCCTTCCTTA CCTGAAGTTG 180  
 CAGGTAGGCC CTGGGTNAGG GCGNGGGGCG CTGGACANTN TCTGGNCCTG GCTGCCCGCT 240  
 3 241

## (2) INFORMATION FOR SEQ ID NO:23:

- (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 497 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:23:

GGCAGAGCA GGGTCTGTN TCCGCCCTGA GCCGCGCTCT NCCTGCTCCA GCAAGGACCA 60

TGAGGGCGCT GGAGGGGCCA GGCCTGTCGC TGCTGTCCTG GTGTTGGCGC TGCTGCCCCT	120
GCTGCCGGTG CCGGCTGTAC GCGGAGTGGC AGAAACACNN ACNTACCCCT GCGGGGACGN	180
AGAGACAGGG GAGCGGCTGG TGTNTNCCCA NTGCCCCCAG GCACCTTTNT GCAGCGGCCG	240
TGCCGNCGAG ACAGCCCCAC GACGTGTGGC CCGTNTCCAC CGCGCCACTA CACGCATTCT	300
GGAACCTACCT GGAGCGCTGN CCTTACTNCA ACGTCCTCTG CGGGGAGCGT NAGGAGGAGG	360
CACGGGTTTN CCACGNCAAC CACAACCCNG GNTTACCGTN GCCGNACCGG TTTCTTCGNG	420
GCAAGTTGGT TTTTNTTTG GAGNAAGGAT TCGTGTNCA ATTNATTJAC GNAGTGATTN	480
NNCNCGGGAA ACTNAAA	497

## (2) INFORMATION FOR SEQ ID NO:24:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 190 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:24:

CGCAACTGCA CGGCCCTGGG ACTGGCCCTC AATGTGCCAG GNTCTTCCTC CCATGACACC	60
CTGTGCACCA GCTGCACTGG CTTCCCCCTC AGACCAGGGT ACCANGAGCT GAGGAGTGTG	120
AGCNTGCCGT CATCGACTTT TTGGCTTTCC AGGACATCTC CATCAAGAGG CTGCAGCGGC	180
TGCTCANGCC	190

## (2) INFORMATION FOR SEQ ID NO:25:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 27 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:25:

CGCGGATCCA CCACTGCCCG GCAGGAG



## (2) INFORMATION FOR SEQ ID NO:26:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 30 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (genomic)

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:26:

GCGTCTAGAC TAGTAATGAG AAGAGGCAGG

30

## (2) INFORMATION FOR SEQ ID NO:27:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 26 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (genomic)

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:27:

CGCCCATGGC AGAAACACCC ACCTAC

26

## (2) INFORMATION FOR SEQ ID NO:28:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 26 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (genomic)

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:28:

CGCAAGCTTC TCTTTCAGTG CAAGTG

26

## (2) INFORMATION FOR SEQ ID NO:29:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 28 base pairs

- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:29:

CGCAAGCTTC TCCTCAGCTC CTGCAGTG

28

(2) INFORMATION FOR SEQ ID NO:30:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 35 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:30:

CGCTCTAGAC CGCCATCATG GCCCGGATCC CCAAG

35

(2) INFORMATION FOR SEQ ID NO:31:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 36 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:31:

CGCGGATCCG CCATCATGAG GCGTGGAGG GGCCAG

36

(2) INFORMATION FOR SEQ ID NO:32:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 30 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:32:

GCGTCTAGAC TAGTAATGAG AAGAGGCAGG

30

(2) INFORMATION FOR SEQ ID NO:33:

- (i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 26 base pairs  
(B) TYPE: nucleic acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:33:

CGCGGTACCC TCTTTCAGTG CAAGTG

26

(2) INFORMATION FOR SEQ ID NO:34:

- (i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 28 base pairs  
(B) TYPE: nucleic acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:34:

CGCGGTACCC TCCTCAGCTC CTGCAAGTG

28

(2) INFORMATION FOR SEQ ID NO:35:

- (i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 34 base pairs  
(B) TYPE: nucleic acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:35:

CGCGAATTCC GCCATCATGG CCCGGATCCC CAAG

34

(2) INFORMATION FOR SEQ ID NO:36:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 27 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:36:

GCGTCTAGAG TAATGAGAAG AGGCAGG

27

(2) INFORMATION FOR SEQ ID NO:37:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 35 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:37:

CGCTCTAGAC CGCCATCATG GCCCGGATCC CCAAG

35

(2) INFORMATION FOR SEQ ID NO:38:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 30 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:38:

CGCTCTAGAC TAGTAATGAG AAGAGGCAGG

30

**What Is Claimed Is:**

1. An isolated nucleic acid molecule comprising a polynucleotide having a nucleotide sequence at least 95% identical to a sequence selected from the group consisting of:

(a) a nucleotide sequence encoding a TNFR polypeptide having the complete amino acid sequence in SEQ ID NO:2, 4 or 6 or as encoded by a cDNA clone contained in ATCC Deposit No. 97798, 97810 or 97809;

(b) a nucleotide sequence encoding a mature TNFR polypeptide having an amino acid sequence at positions 27-259, 31-300 or 31-170 in SEQ ID NO:2, 4 and 6, respectively, or as encoded by the cDNA clone contained in the ATCC Deposit No. 97798, 97810 or 97809;

(c) a nucleotide sequence encoding the soluble extracellular domain of a TNFR polypeptide having the amino acid sequence at positions 27-240, 31-283 or 31-166 of SEQ ID NOS:2, 4 and 6, respectively; and

(d) a nucleotide sequence complementary to any of the nucleotide sequences in (a), (b), or (c) above.

2. The nucleic acid molecule of claim 1 wherein said polynucleotide has a complete nucleotide sequence selected from the group consisting of SEQ ID NO:1, SEQ ID NO:3 and SEQ ID NO:5.

3. The nucleic acid molecule of claim 1 wherein said polynucleotide has a nucleotide sequence which encodes a TNFR polypeptide having a complete amino acid sequence selected from the group consisting of SEQ ID NO:2, SEQ ID NO:4 and SEQ ID NO:6.

4. The nucleic acid molecule of claim 1 wherein said polynucleotide has a nucleotide sequence encoding the mature form of a TNFR polypeptide having an amino acid sequence from about 27 to about 259 in SEQ ID NO:2, from about 31 to about 300 in SEQ ID NO:4, or from about 31 to about 170 in SEQ ID NO:6.

5. The nucleic acid molecule of claim 1 wherein said polynucleotide has a nucleotide sequence encoding the soluble extracellular domain of a TNFR polypeptide having the amino acid sequence from about 27 to about 240 in SEQ ID NO:2, from about 31 to about 283 in SEQ ID NO:4, or from about 31 to about 166 of SEQ ID NO:6.

6. An isolated nucleic acid molecule comprising a polynucleotide having a nucleotide sequence at least 95% identical to a sequence selected from the group consisting of:

- (a) a nucleotide sequence encoding a polypeptide comprising the amino acid sequence of residues m-259 of SEQ ID NO:2, where m is an integer in the range of 1-53;
- (b) a nucleotide sequence encoding a polypeptide comprising the amino acid sequence of residues n-300 of SEQ ID NO:4, where n is an integer in the range of 1-49;
- (c) a nucleotide sequence encoding a polypeptide comprising the amino acid sequence of residues n-170 of SEQ ID NO:6, where n is an integer in the range of 1-49;
- (d) a nucleotide sequence encoding a polypeptide comprising the amino acid sequence of residues 1-x of SEQ ID NO:2, where x is an integer in the range of 149-259;
- (e) a nucleotide sequence encoding a polypeptide comprising the amino acid sequence of residues 1-y of SEQ ID NO:4, where y is an integer in the range of 193-300;
- (f) a nucleotide sequence encoding a polypeptide comprising the amino acid sequence of residues 1-z of SEQ ID NO:6, where z is an integer in the range of 132-170; and
- (g) a nucleotide sequence encoding a polypeptide having the amino acid sequence consisting of residues m-x of SEQ ID NO:2, n-y of SEQ ID NO:4, or n-z of SEQ ID NO:6, where m, n, x, y and z are defined in (a), (b), (c), (d), (e) and (f) above.

7. An isolated nucleic acid molecule comprising a polynucleotide having a nucleotide sequence at least 95% identical to a sequence selected from the group consisting of:

- (a) a nucleotide sequence encoding a polypeptide consisting of a portion of a complete TNFR amino acid sequence encoded by a cDNA clone contained in ATCC Deposit No. 97798, 97810 or 97809 wherein said portion excludes from 1 to about 52, from 1 to about 48 and from 1 to about 48 amino acids from the amino terminus of said complete amino acid sequence encoded by the cDNA clone contained in ATCC Deposit No. 97798, 97810 and 97809, respectively;

(b) a nucleotide sequence encoding a polypeptide consisting of a portion of a complete TNFR amino acid sequence encoded by a cDNA clone contained in ATCC Deposit No. 97798, 97810 or 97809 wherein said portion excludes from 1 to about 110, from 1 to about 107, or from 1 to about 38 amino acids from the carboxy terminus of said complete amino acid sequence encoded by the cDNA clone contained in ATCC Deposit No. 97798, 97810 and 97809, respectively; and

(c) a nucleotide sequence encoding a polypeptide consisting of a portion of a complete TNFR amino acid sequence encoded by the cDNA clone contained in ATCC Deposit No. 97798, 97810 or 97809, wherein said portion includes a combination of any of the amino terminal and carboxy terminal deletions for the respective clones in (a) and (b), above.

8. The nucleic acid molecule of claim 1 wherein said polynucleotide has the complete nucleotide sequence of the cDNA clone contained in ATCC Deposit No. 97798, 97810 or 97809.

9. The nucleic acid molecule of claim 1 wherein said polynucleotide has the nucleotide sequence encoding a TNFR polypeptide having the complete amino acid sequence encoded by the cDNA clone contained in ATCC Deposit No. 97798, 97810 or 97809.

10. The nucleic acid molecule of claim 1 wherein said polynucleotide has the nucleotide sequence encoding a mature TNFR polypeptide having the amino acid sequence encoded by the cDNA clone contained in ATCC Deposit No. 97798, 97810 or 97809.

11. An isolated nucleic acid molecule comprising a polynucleotide which hybridizes under stringent hybridization conditions to a polynucleotide having a nucleotide sequence identical to a nucleotide sequence in (a), (b), (c), or (d) of claim 1 wherein said polynucleotide which hybridizes does not hybridize under stringent hybridization conditions to a polynucleotide having a nucleotide sequence consisting of only A residues or of only T residues.

12. An isolated nucleic acid molecule comprising a polynucleotide which encodes the amino acid sequence of an epitope-bearing portion of a TNFR polypeptide having an amino acid sequence in (a), (b), (c) or (d) of claim 1.

13. The isolated nucleic acid molecule of claim 12, which encodes an epitope-bearing portion of a TNFR polypeptide comprising amino acid residues selected from the group consisting of: from about Gln-42 to about Glu-52 in SEQ ID NO:2, from about His-58 to about Cys-66 in SEQ ID NO:2, from about Pro-68 to about Thr-76 in SEQ ID NO:2, from about Ser-79 to about Cys-85 in SEQ ID NO:2, from about Cys-91 to about Thr-102 in SEQ ID NO:2, from about Gln-110 to about Pro-122 in SEQ ID NO:2, from about Arg-126 to about Val-136 in SEQ ID NO:2, from about Thr-142 to about Glu-148 in SEQ ID NO:2, from about Ala-31 to about Thr-46 in SEQ ID NO:4, from about Phe-57 to about Thr-117 in SEQ ID NO:4, from about Cys-132 to about Thr-175 in SEQ ID NO:4, from about Gly-185 to about Thr-194 in SEQ ID NO:4, from about Val-205 to about Asp-217 in SEQ ID NO:4, from about Pro-239 to about Leu-264 in SEQ ID NO:4, and from about Ala-283 to about Pro-298 in SEQ ID NO:4, from about Ala-31 to about Thr-46 in SEQ ID NO:6, from about Phe-57 to about Gln-80 in SEQ ID NO:6, from about Glu-86 to about His-106 in SEQ ID NO:6, from about Thr-108 to about Phe-119 in SEQ ID NO:6, from about His-129 to about Val-138 in SEQ ID NO:6, and from about Gly-142 to about Pro-166 in SEQ ID NO:6 to about in SEQ ID NO:6.

14. A method for making a recombinant vector comprising inserting an isolated nucleic acid molecule of claim 1 into a vector.

15. A recombinant vector produced by the method of claim 14.

16. A method of making a recombinant host cell comprising introducing the recombinant vector of claim 15 into a host cell.

17. A recombinant host cell produced by the method of claim 16.



18. A recombinant method for producing a TNFR polypeptide, comprising culturing the recombinant host cell of claim 17 under conditions such that said polypeptide is expressed and recovering said polypeptide.

19. An isolated TNFR polypeptide comprising an amino acid sequence at least 95% identical to a sequence selected from the group consisting of:

(a) the amino acid sequence of a full-length TNFR polypeptide having the complete amino acid sequence shown in SEQ ID NO:2, 4 or 6, or as encoded by a cDNA clone contained in ATCC Deposit No. 97798, 97810 or 97809;

(b) the amino acid sequence of a mature TNFR polypeptide having the amino acid sequence at positions 27-259 in SEQ ID NO:2, 31-300 in SEQ ID NO:4, or 31-170 in SEQ ID NO:6, or as encoded by a cDNA clone contained in ATCC Deposit No. 97798, 97810 or 97809; or

(c) the amino acid sequence of a soluble extracellular domain of a TNFR polypeptide having the amino acid sequence at positions 27 to 240 in SEQ ID NO:2, 31 to 283 in SEQ ID NO:4, or 31 to 166 in SEQ ID NO:6, or as encoded by the cDNA clone contained in ATCC Deposit No. 97798, 97810 or 97809.

20. An isolated polypeptide comprising an epitope-bearing portion of the TNFR protein, wherein said portion is selected from the group consisting of a polypeptide comprising amino acid residues: from about Gln-42 to about Glu-52 in SEQ ID NO:2, from about His-58 to about Cys-66 in SEQ ID NO:2, from about Pro-68 to about Thr-76 in SEQ ID NO:2, from about Ser-79 to about Cys-85 in SEQ ID NO:2, from about Cys-91 to about Thr-102 in SEQ ID NO:2, from about Gln-110 to about Pro-122 in SEQ ID NO:2, from about Arg-126 to about Val-136 in SEQ ID NO:2, from about Thr-142 to about Glu-148 in SEQ ID NO:2, from about Ala-31 to about Thr-46 in SEQ ID NO:4, from about Phe-57 to about Thr-117 in SEQ ID NO:4, from about Cys-132 to about Thr-175 in SEQ ID NO:4, from about Gly-185 to about Thr-194 in SEQ ID NO:4, from about Val-205 to about Asp-217 in SEQ ID NO:4, from about Pro-239 to about Leu-264 in SEQ ID NO:4, and from about Ala-283 to about Pro-298 in SEQ ID NO:4, from about Ala-31 to about Thr-46 in SEQ ID NO:6, from about Phe-57 to about Gln-80 in SEQ ID NO:6, from about Glu-86 to about His-106 in SEQ ID NO:6, from about Thr-108 to about Phe-119 in SEQ ID NO:6, from about His-129 to about Val-138 in SEQ ID NO:6, and from about Gly-142 to about Pro-166 in SEQ ID NO:6 to about in SEQ ID NO:6.

21. An isolated antibody that binds specifically to a TNFR polypeptide of claim 19.

22. An isolated nucleic acid molecule comprising a polynucleotide having a sequence at least 95% identical to a sequence selected from the group consisting of:

- (a) the nucleotide sequence of clone HPRCB54R (SEQ ID NO:19);
- (b) the nucleotide sequence of clone HSJAU57RA (SEQ ID NO:20);
- (c) the nucleotide sequence of clone HELBP70R (SEQ ID NO:21);
- (d) the nucleotide sequence of clone HUSCB54R (SEQ ID NO:22);
- (e) the nucleotide sequence of clone HELDI06R (SEQ ID NO:23);
- (f) the nucleotide sequence of clone HCEOW38R (SEQ ID NO:24); and
- (g) a nucleotide sequence complementary to any of the nucleotide sequences in (a), (b), (c), (d), (e), or (f) above.

### *Abstract*

The present invention relates to novel Tumor Necrosis Factor Receptor proteins. In particular, isolated nucleic acid molecules are provided encoding the human TNFR-5, -6 $\alpha$  & -6 $\beta$  proteins. TNFR-5, -6 $\alpha$  & -6 $\beta$  polypeptides are also provided as are vectors, host cells and recombinant methods for producing the same. The invention further relates to screening methods for identifying agonists and antagonists of TNFR-5, -6 $\alpha$  & -6 $\beta$  activity. Also provided are diagnostic methods for detecting immune system-related disorders and therapeutic methods for treating immune system-related disorders.



Figure 1

CCTCTCCACGCGCACGAACTCAGCCAACGATTTCTGATAGATTTTTGGGAGTTTGACCAG  
AGATGCAAGGGGTGAAGGAGCGCTTCTACCGTTAGGGAACCTCTGGGGACAGAGCGCCCC  
GGCCGCCTGATGGCCGAGGCAGGGTGCACCCAGGACCCAGGACGGCGTCGGGAACCATA  
CCATGGGCGGATCCCCAAGACCCTAAAGTTCTGTCGTCGTCATCGTCGCGGTCTGCTGC  
E A R I P K T L K F V V V I V A V L L L P  
CAGTCCTAGCTTACTCTGCCACCACTGCCCCGAGGAGGAAGTTCCCCAGCAGACAGTGG  
V L A Y S A T T A R Q E E V P Q Q T V A  
CCCCACAGCAACAGAGGCACAGCTTCAAGGGGGAGGAGTGTCCAGCAGGATCTCATAGAT  
P Q Q Q R H S F K G E E C P A G S H R S  
CAGAACATACTGGAGCCTGTAACCCGTGCACAGAGGGTGTGGATTACACCAACGCTTCCA  
E H T G A C N P C T E G V D Y T N A S N  
ACAATGAACCTTCTTGCTTCCATGTACAGTTTGTAAATCAGATCAAAAACA1AAAAGTT  
N E P S C F P C T V C K S D Q K H K S S  
CCTGCACCATGACCAGAGACACAGTGTGTCTAGTGTAAAGAAGGCACCTTCCGGAATGAAA  
C T M T R D T V C Q C K E G T F R N E N  
ACTCCCCAGAGATGTGCCGGAAGTGTAGCAGGTGCCCTAGTGGGGAAGTCCAAGTCAGTA  
S P E M C R K C S R C P S G E V Q V S N  
ATTGTACGTCTGGGATGATATCCAGTGTGTTGAAGAATTTGGTGCCAATGCCACTGTGG  
C T S W D D I Q C V E E F G A N A T V E  
AAACCCAGCTGCTGAAGAGACAATGAACACCAGCCCGGGGACTCCTGCCCCAGCTGCTG  
T P A A E E T M N T S P G T P A P A A E  
AAGAGACAATGAACACCAGCCAGGGACTCCTGCCCCAGCTGCTGAAGAGACAATGACCA  
E T M N T S P G T P A P A A E E T M T T  
CCAGCCCGGGGACTCCTGCCCCAGCTGCTGAAGAGACAATGACCACCAGCCCGGGGACTC  
S P G T P A P A A E E T M T T S P G T P  
CTGCCCCAGCTGCTGAAGAGACAATGACCACCAGCCCGGGGACTCCTGCCTCTTCTCATT  
A P A A E E T M T T S P G T P A S S H Y



Figure 1 (continued)

ACCTCTCATGCACCATCGTAGGGATCATAGTTCTAATTGTGCTTCTGATTGTGTTTGT  
L S C T I V G I I V L I V L L I V F V \*  
GAAAGACTTCACTGTGGAAGAAATTCCTTCCTTACCTGAAAGGTTACAGGTAGGCGCTGGC  
TGAGGGCGGGGGGGCGCTGGACACTCTCTGCCCTGCCTCCCTCTGCTGTGTGCCACAGAC  
AGAAACGCCTGCCCCGCCCCAAGTCCTGGTGCTCCAGCCTGGCTCTATCTTCCTCCTT  
GTGATCGTCCCATCCCCACATCCCGTGACCCCCCAGGACCCTGGTCTCATCAGTCCCTC  
TCCTGGAGCTGGGGGTCCACACATCTCCAGCCAAGTCCAAGAGGCAGGGCCAGTTCCTC  
CCATCTTCAGGCCAGCCAGGCAGGGGGCAGTCGGCTCCTCAACTGGGTGACAAGGGTGA  
GGATGAGAAGTGGTCACGGGATTATTTCAGCCTTGGTCAGAGCAGAACACAGAGATTTTC  
CGTGAAAAAAAA



Figure 2

GCTCTCCCTGCTCCAGCAAGGACCATGAGGGCGCTGGAGGGGCCAGGCGTGTGCTGCTG  
M R A L E G P G L S L L

TGCCTGGTGTGGCGCTGCCCTGCTGCCGGTGCCGGTGTACGCGGAGTGGCAGAA  
C L V L A L P A L L P V P A V R G V A E

ACACCCACCTACCCCTGGCGGGACGCAGAGACAGGGGAGCGGCTGGTGTGCGCCAGTGC  
T P T Y P W R D A E T G E R L V C A Q C

CCCCCAGGCACCTTGTGTCAGCGCGCTGCCGCCGAGACAGCCCCACGACGTGTGGCCCC  
P P G T F V Q R P C R R D S P T T C G P

TGTCCACCGCGCCACTACACGCAGTTCTGGAACCTGGAGCGCTGCCGCTACTGCAAC  
C P P R H Y T Q F W N Y L E R C R Y C N

GTCCTCTGCGGGGAGCGTGAGGAGGAGGCACGGGCTTGCCACGCCACCCACAACCGTGCC  
V L C G E R E E E A R A C H A T H N R A

TGCCGCTGCCGCACCGCTTCTTCGCGCACGCTGGTTTCTGCTTGGAGCACGCATCGTGT  
C R C R T G F F A H A G F C L E H A S C

CCACCTGGTGCGGGCGTGATTGCCCGGGCACCCCCAGCCAGAACACGCAGTGCCAGCCG  
P P G A G V I A P G T P S Q N T Q C Q P

TGCCCCCAGGCACCTTCTCAGCCAGCAGCTCCAGCTCAGAGCAGTGCCAGCCCCACCGC  
C P P G T F S A S S S S S E Q C Q P H R

AACTGCACGGCCCTGGGCCTGGCCCTCAATGTGCCAGGCTCTTCTCCCATGACACCTG  
N C T A L G L A L N V P G S S S H D T L

TGCACCAGCTGCACTGGCTTCCCCCTCAGCACCAGGTACCAGGAGCTGAGGAGTGTGAG  
C T S C T G F P L S T R V P G A E E C E

CGTGCCGTCATCGACTTTGTGGCTTCCAGGACATCTCCATCAAGAGGCTGCAGCGGCTG  
R A V I D F V A F Q D S I K R L Q R L

CTGCAGGCCCTCGAGGCCCCGAGGGCTGGGGTCCGACACCAAGGGCGGGCCGCGCGCC  
L Q A L E A P E G W G P T P R A G R A A

TTGCAGCTGAAGCTGCGTGGCGGCTCACGGAGCTCCTGGGGCGCAGGACGGGGCGCTG  
L Q L K L R R R L T E L L G A Q D G A L

CTGGTGGCGCTGCTGCAGGCGCTGCGCGTGGCCAGGATGCCCGGGCTGGAGCGGAGCGTC  
L V R L L Q A L R V A R M P G L E R S V

CGTGAGCGCTTCTCCCTGTGCACTGATCCTGGCCCCCTCTATTATTCTACATCCTTG  
R E R F L P V H \*

GCACCCCACTTGCACTGAAAGAGGCTTTTTTTTAAATAGAAGAAATGAGGTTTCTTAAAG  
CTTATTTTATAAAGCTTTTTCATAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAA

Figure 3

TGGCATGTCCGGTCAGGCACAGCAGGGTCCTGTGTCCGGCTGAGCCGGCTCTCCCTGCT  
CCAGCAAGGACCATGAGGGCGCTGGAGGGGCCAGGCCTGTGCTGCTGTGCCTGGTGTG  
M R A L E G P G L S L L C L V L  
GCCGTGCCTGCCCTGTGCGGGTGCCGGCTGTACGGGAGTGGCAGAAACACCCACCTAC  
A L P A L L P V P A V R G V A E T P T Y  
CCCTGGCGGGACGCAGAGACAGGGGAGCGGCTGGTGTGCGCCAGTGCCCCCAGGCACC  
P W R D A E T G E R L V C A Q C P P G T  
TTTGTGCAGCGGCGTGCCGCCGAGACAGCCCCACGACGTGTGGCCCGTGTCCACCGCGC  
F V Q R P C R R D S P T T C G P C P P R  
CACTACACGCAGTTCTGGAACCTACCTGGAGCGCTGCCGCTACTGCAACGTCTCTGCGGG  
I Y T Q F W N Y L E R C R Y C N V L C G  
GAGCGTGAGGAGGAGGCACGGGCTTGCCACGCCACCCACAACCGTGCTGCCGCTGCCCC  
E R E E E A R A C H A T H N R A C R C R  
ACCGGCTTCTTCGCGCAGCGTGGTTTCTGCTTGGAGCAGCATCGTGTCCACCTGGTGGC  
T G F F A H A G F C L E H A S C P P G A  
GGCGTGATTGCCCGGGTGAGAGCTGGGCGAGGGGAGGGGCCCCCAGGAGTGGTGGCCGG  
G V I A P G E S W A R G G A P R S G G R  
AGGTGTGGCAGGGGTGAGGTGGTGGTCCAGCCTTGCAACCTGAGCTAGGACACCAAGT  
F C G R G Q V A G P S L A P \*  
CCCTGACCTGTTCTTCCCTCCTGGCTGCAGGCACCCCCAGCCAGAACACGCAGTGCCA  
GCGTGCCCCCAGGCACCTTCTCAGCCAGCAGCTCCAGCTCAGAGCACTGCCAGCCCA  
CCGCAACTGCAGGCCCCCTGGCCTGGCCCTCAATGTGCCAGGCTCTTCTCCCATGACAC  
CCTGTGCACCAGCTGCACTGGCTTCCCCCTCAACACAGGGTACCAGGTGAGCCAGAGGC  
CTGAGGGGGCAGTACACTGCAGGCCAGGCCCACTTGTGCCCTCACTCCTGCCCTGCACG  
TGCACTAGCCTGAGGCATGCCAGCTGGCTCTGGGAAGGGGCCACAGTGGATTGAGGGG  
TACAGGGCTCCCTCCACTAGATCCCCACCAAGTCTGCCCTCTCAGGGGTGGCTGAGAATTT  
GGATCTGAGCCAGGGCACAGCCTCCCTGGAGAGCTCTGGGAAAGTGGGCAGCAATCTCC

Figure 3 (continued)

TAACTGCCCCGAGGGGAAGGTGGCTGGCTCCTCTGACACGGGGAACCGAGGCCTGATGGT  
AACTCTCCTAACTGCCTGAGAGGAAGGTGGCTGCCCTCCTCTGACATGGGAAACCGAGGC  
CCAATGTTAACCCTGTTGAGAAGTCACAGSGGGAAGTACCCCCCTTAACATCAAGTCAG  
GTCCGGTCCATCTGCAGGTCCCAACTCGCCCCCTTCGATGGCCCAGGAGCCCAAGCCCT  
TGCTGGGCCCCCTTGCTCTTGAGCCAAGGTCCGAGTGGCCGCTCCTGCCCCCTAGGC  
CTTGCTCCAGCTCTCTGACCGAAGGCTCCTGCCCCCTTCTCCAGTCCCCATCGTTGCACT  
GCCCTCTCCAGCACGGCTCACTGCACAGGGATTCTCTCTCCTGCAACCCCCCGAGTGG  
GGGACAGAAAGCAGGGTACCTGGCAGCCCCCGCCAGTGTGTGGGTGAAATGATCGGAC  
CGCTGCCTCCCCACCCCACTGCAGGAGCTGAGGAGTGTGAGCGTGCCGTCATCGACTTTG  
TCCTTTCCAGGACATCTCCATCAAGAGGAGCGGCTGCTGCAGGCC



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Figure 4





Figure 4 (continued)

194	.....IENVKGTDSSTTVLLPLIFFG-L.....	TFR1
212	GAVHLCVYSTNSONTOTPEPSTAPSTLPLMGPSPPA	TFR2
220	TPTEGSDSTAPSTOEPEAPEDLIASL.....VAGVTTV	NFR
224	LAILC.....LNR	
228	.....LTSNTRCKEGRSRL.....GML.....PNS	
230	.....ARSGAUSPHPOPHLPVSEMLEAR.....TAGHMTL	0127
234	AASKLTRADSPSSVGRPSDPLSPTOP.....CPEGSGDCR	0100
238	ALVVIP.....CNO	
242	DLSPGASSVTPAPAREPGHPOIIS.....4-1B8	
246	.....RDPATOPDETGGP.....	0N40
250	GL-ESILT.....SELTITMNTDCN	V22
254	GL-ESIST.....SELTITMNRKDCD	0N8
258	.....FG.....TFR-11a	
262	.....CTGF.....PLSTRVPG	TFR-6a
266	.....TFR-6b	
270	.....TFR-6b	
274	.....TFR-6b	
278	.....TFR-6b	
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1082	.....TFR-6b	
1086	.....TFR-6b	
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1094	.....TFR-6b	
1098	.....TFR-6b	
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1106	.....TFR-6b	
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1186	.....TFR-6b	
1190	.....TFR-6b	
1194	.....TFR-6b	
1198	.....TFR-6b	
1202	.....TFR-6b	
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1642	.....TFR-6b	
1646	.....TFR-6b	
1650	.....TFR-6b	
1654	.....TFR-6b	

Figure 4 (continued)

204	TRPLAPNPSFS	TPGFT	TLGFSPT	SSTFTSSSTVTPGD	TRU
205	LOP	VPMLA	DRAR	TQGEQOHLITA	TRQ
206	NKOGASRQV	NOTPP	EGENLNSDGSVD	NFR	
207	PEGESPPCA	PRADPHFDLAEP	LL	LIR	
208	SPTLNE	TVAINLSVDLSRYIT		PNS	
209	TFEAPLGC	DCPT	NCEAPASTSPTOSLLVDSQA	CD30	
210	PNKAPHKCL	PO	EINFPD	CD40	
211	SOGPSTRV	E	VGGRAVAAILGLCL	OM40	
212	LNFIICN	K	S-SFKO	LTK	VZ2
213	LNFIICN	K	DSYSSKO	LTK	OM8
214	TSPGTPAPAAETM	TS	GTPAPAAETMTTSPCT		TRR-11be
215	LOLLLOALEA	CM	SA		TRR-6a
216	LOLLLOALEA	CM	SA		TRR-6b
217	LOLLLOALEA	CM	SA		TRR-6c
218	LOLLLOALEA	CM	SA		TRR-6d
219	LOLLLOALEA	CM	SA		TRR-6e
220	LOLLLOALEA	CM	SA		TRR-6f
221	LOLLLOALEA	CM	SA		TRR-6g
222	LOLLLOALEA	CM	SA		TRR-6h
223	LOLLLOALEA	CM	SA		TRR-6i
224	LOLLLOALEA	CM	SA		TRR-6j
225	LOLLLOALEA	CM	SA		TRR-6k
226	LOLLLOALEA	CM	SA		TRR-6l
227	LOLLLOALEA	CM	SA		TRR-6m
228	LOLLLOALEA	CM	SA		TRR-6n
229	LOLLLOALEA	CM	SA		TRR-6o
230	LOLLLOALEA	CM	SA		TRR-6p
231	LOLLLOALEA	CM	SA		TRR-6q
232	LOLLLOALEA	CM	SA		TRR-6r
233	LOLLLOALEA	CM	SA		TRR-6s
234	LOLLLOALEA	CM	SA		TRR-6t
235	LOLLLOALEA	CM	SA		TRR-6u
236	LOLLLOALEA	CM	SA		TRR-6v
237	LOLLLOALEA	CM	SA		TRR-6w
238	LOLLLOALEA	CM	SA		TRR-6x
239	LOLLLOALEA	CM	SA		TRR-6y
240	LOLLLOALEA	CM	SA		TRR-6z
241	LOLLLOALEA	CM	SA		TRR-6aa
242	LOLLLOALEA	CM	SA		TRR-6ab
243	LOLLLOALEA	CM	SA		TRR-6ac
244	LOLLLOALEA	CM	SA		TRR-6ad
245	LOLLLOALEA	CM	SA		TRR-6ae
246	LOLLLOALEA	CM	SA		TRR-6af
247	LOLLLOALEA	CM	SA		TRR-6ag
248	LOLLLOALEA	CM	SA		TRR-6ah
249	LOLLLOALEA	CM	SA		TRR-6ai
250	LOLLLOALEA	CM	SA		TRR-6aj
251	LOLLLOALEA	CM	SA		TRR-6ak
252	LOLLLOALEA	CM	SA		TRR-6al
253	LOLLLOALEA	CM	SA		TRR-6am
254	LOLLLOALEA	CM	SA		TRR-6an
255	LOLLLOALEA	CM	SA		TRR-6ao
256	LOLLLOALEA	CM	SA		TRR-6ap
257	LOLLLOALEA	CM	SA		TRR-6aq
258	LOLLLOALEA	CM	SA		TRR-6ar
259	LOLLLOALEA	CM	SA		TRR-6as
260	LOLLLOALEA	CM	SA		TRR-6at
261	LOLLLOALEA	CM	SA		TRR-6au
262	LOLLLOALEA	CM	SA		TRR-6av
263	LOLLLOALEA	CM	SA		TRR-6aw
264	LOLLLOALEA	CM	SA		TRR-6ax
265	LOLLLOALEA	CM	SA		TRR-6ay
266	LOLLLOALEA	CM	SA		TRR-6az
267	LOLLLOALEA	CM	SA		TRR-6ba
268	LOLLLOALEA	CM	SA		TRR-6bb
269	LOLLLOALEA	CM	SA		TRR-6bc
270	LOLLLOALEA	CM	SA		TRR-6bd
271	LOLLLOALEA	CM	SA		TRR-6be
272	LOLLLOALEA	CM	SA		TRR-6bf
273	LOLLLOALEA	CM	SA		TRR-6bg
274	LOLLLOALEA	CM	SA		TRR-6bh
275	LOLLLOALEA	CM	SA		TRR-6bi
276	LOLLLOALEA	CM	SA		TRR-6bj
277	LOLLLOALEA	CM	SA		TRR-6bk
278	LOLLLOALEA	CM	SA		TRR-6bl
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284	LOLLLOALEA	CM	SA		TRR-6br
285	LOLLLOALEA	CM	SA		TRR-6bs
286	LOLLLOALEA	CM	SA		TRR-6bt
287	LOLLLOALEA	CM	SA		TRR-6bu
288	LOLLLOALEA	CM	SA		TRR-6bv
289	LOLLLOALEA	CM	SA		TRR-6bw
290	LOLLLOALEA	CM	SA		TRR-6bx
291	LOLLLOALEA	CM	SA		TRR-6by
292	LOLLLOALEA	CM	SA		TRR-6bz
293	LOLLLOALEA	CM	SA		TRR-6ca
294	LOLLLOALEA	CM	SA		TRR-6cb
295	LOLLLOALEA	CM	SA		TRR-6cc
296	LOLLLOALEA	CM	SA		TRR-6cd
297	LOLLLOALEA	CM	SA		TRR-6ce
298	LOLLLOALEA	CM	SA		TRR-6cf
299	LOLLLOALEA	CM	SA		TRR-6cg
300	LOLLLOALEA	CM	SA		TRR-6ch
301	LOLLLOALEA	CM	SA		TRR-6ci
302	LOLLLOALEA	CM	SA		TRR-6cj
303	LOLLLOALEA	CM	SA		TRR-6ck
304	LOLLLOALEA	CM	SA		TRR-6cl
305	LOLLLOALEA	CM	SA		TRR-6cm
306	LOLLLOALEA	CM	SA		TRR-6cn
307	LOLLLOALEA	CM	SA		TRR-6co
308	LOLLLOALEA	CM	SA		TRR-6cp
309	LOLLLOALEA	CM	SA		TRR-6cq
310	LOLLLOALEA	CM	SA		TRR-6cr
311	LOLLLOALEA	CM	SA		TRR-6cs
312	LOLLLOALEA	CM	SA		TRR-6ct
313	LOLLLOALEA	CM	SA		TRR-6cu
314	LOLLLOALEA	CM	SA		TRR-6cv
315	LOLLLOALEA	CM	SA		TRR-6cw
316	LOLLLOALEA	CM	SA		TRR-6cx
317	LOLLLOALEA	CM	SA		TRR-6cy
318	LOLLLOALEA	CM	SA		TRR-6cz
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325	LOLLLOALEA	CM	SA		TRR-6dg
326	LOLLLOALEA	CM	SA		TRR-6dh
327	LOLLLOALEA	CM	SA		TRR-6di
328	LOLLLOALEA	CM	SA		TRR-6dj
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335	LOLLLOALEA	CM	SA		TRR-6dq
336	LOLLLOALEA	CM	SA		TRR-6dr
337	LOLLLOALEA	CM	SA		TRR-6ds
338	LOLLLOALEA	CM	SA		TRR-6dt
339	LOLLLOALEA	CM	SA		TRR-6du
340	LOLLLOALEA	CM	SA		TRR-6dv
341	LOLLLOALEA	CM	SA		TRR-6dw
342	LOLLLOALEA	CM	SA		TRR-6dx
343	LOLLLOALEA	CM	SA		TRR-6dy
344	LOLLLOALEA	CM	SA		TRR-6dz
345	LOLLLOALEA	CM	SA		TRR-6ea
346	LOLLLOALEA	CM	SA		TRR-6eb
347	LOLLLOALEA	CM	SA		TRR-6ec
348	LOLLLOALEA	CM	SA		TRR-6ed
349	LOLLLOALEA	CM	SA		TRR-6ee
350	LOLLLOALEA	CM	SA		TRR-6ef
351	LOLLLOALEA	CM	SA		TRR-6eg
352	LOLLLOALEA	CM	SA		TRR-6eh
353	LOLLLOALEA	CM	SA		TRR-6ei
354	LOLLLOALEA	CM	SA		TRR-6ej
355	LOLLLOALEA	CM	SA		TRR-6ek
356	LOLLLOALEA	CM	SA		TRR-6el
357	LOLLLOALEA	CM	SA		TRR-6em
358	LOLLLOALEA	CM	SA		TRR-6en
359	LOLLLOALEA	CM	SA		TRR-6eo
360	LOLLLOALEA	CM	SA		TRR-6ep
361	LOLLLOALEA	CM	SA		TRR-6eq
362	LOLLLOALEA	CM	SA		TRR-6er
363	LOLLLOALEA	CM	SA		TRR-6es
364	LOLLLOALEA	CM	SA		TRR-6et
365	LOLLLOALEA	CM	SA		TRR-6eu
366	LOLLLOALEA	CM	SA		TRR-6ev
367	LOLLLOALEA	CM	SA		TRR-6ew
368	LOLLLOALEA	CM	SA		TRR-6ex
369	LOLLLOALEA	CM	SA		TRR-6ey
370	LOLLLOALEA	CM	SA		TRR-6ez
371	LOLLLOALEA	CM	SA		TRR-6fa
372	LOLLLOALEA	CM	SA		TRR-6fb
373	LOLLLOALEA	CM	SA		TRR-6fc
374	LOLLLOALEA	CM	SA		TRR-6fd
375	LOLLLOALEA	CM	SA		TRR-6fe
376	LOLLLOALEA	CM	SA		TRR-6ff
377	LOLLLOALEA	CM	SA		TRR-6fg
378	LOLLLOALEA	CM	SA		TRR-6fh
379	LOLLLOALEA	CM	SA		TRR-6fi
380	LOLLLOALEA	CM	SA		TRR-6fj
381	LOLLLOALEA	CM	SA		TRR-6fk
382	LOLLLOALEA	CM	SA		TRR-6fl
383	LOLLLOALEA	CM	SA		TRR-6fm
384	LOLLLOALEA	CM	SA		TRR-6fn
385	LOLLLOALEA	CM	SA		TRR-6fo
386	LOLLLOALEA	CM	SA		TRR-6fp
387	LOLLLOALEA	CM	SA		TRR-6fq
388	LOLLLOALEA	CM	SA		TRR-6fr
389	LOLLLOALEA	CM	SA		TRR-6fs
390	LOLLLOALEA	CM	SA		TRR-6ft
391	LOLLLOALEA	CM	SA		TRR-6fu
392	LOLLLOALEA	CM	SA		TRR-6fv
393	LOLLLOALEA	CM	SA		TRR-6fw
394	LOLLLOALEA	CM	SA		TRR-6fx
395	LOLLLOALEA	CM	SA		TRR-6fy
396	LOLLLOALEA	CM	SA		TRR-6fz
397	LOLLLOALEA	CM	SA		TRR-6ga
398	LOLLLOALEA	CM	SA		TRR-6gb
399	LOLLLOALEA	CM	SA		TRR-6gc
400	LOLLLOALEA	CM	SA		TRR-6gd
401	LOLLLOALEA	CM	SA		TRR-6ge
402	LOLLLOALEA	CM	SA		TRR-6gf
403	LOLLLOALEA	CM	SA		TRR-6gg
404	LOLLLOALEA	CM	SA		TRR-6gh
405	LOLLLOALEA	CM	SA		TRR-6gi
406	LOLLLOALEA	CM	SA		TRR-6gj
407	LOLLLOALEA	CM	SA		TRR-6gk
408	LOLLLOALEA	CM	SA		TRR-6gl
409	LOLLLOALEA	CM	SA		TRR-6gm
410	LOLLLOALEA	CM	SA		TRR-6gn
411	LOLLLOALEA	CM	SA		TRR-6go
412	LOLLLOALEA	CM	SA		TRR-6gp
413	LOLLLOALEA	CM	SA		TRR-6gq
414	LOLLLOALEA	CM	SA		TRR-6gr
415	LOLLLOALEA	CM	SA		TRR-6gs
416	LOLLLOALEA	CM	SA		TRR-6gt
417	LOLLLOALEA	CM	SA		TRR-6gu

Figure 4 (continued)

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355 NVPPLRMKEFVRFLLSDDNEIDRLLEONGRCLKEACYSHL TNR1
372 PGCHGTOVNVTC VINCSSSDH SSQCSQASSTMGFT TNR2
355 AGDTMHLAELGYOBEHIDFT-EACPVN TNR
377 PGHGOVANHANGINHTGCVTVTCNVIYNGPVLGC LBR
216 KGFVRKMGHAKIDEIKNDNVODTAEQKOLL PMS
214 RRYRSMKGESPVPAEPCRYSCPR COT
441 RRSSTOLRSASCTEPAEERGLMSQPLMETCHSVGAAYL CMO
256 HCCQVVT CMO
222 4-LB
245 ALPPD-AHKPPGGSF CMO
308 AQDYETDTISYRVGNVLODSHMPGSCMINK VC2
311 TODYETDTISYHVGHVLVDHMPGRCDTNK CMB
218 TNR-11b
262 TNR-6a
260 TNR-6a
155 TNR-6b

405 ATMRRTPREATLELLGNVLRDMDLGC TNR1
409 DSSPSES-PKDE--QVPFSKECAF TNR2
388 ASWATQDSTLDA-- TNR
365 RGPGRP-PAPP--EPYPTPECA LBR
279 RNMHQLMKKEA-YDILKDLKKNL--CTLAERIO PMS
239 EECSTI COT
401 ESLPLODASPAGG-PSSPHDLPEPRVSTENKIEKIYI CMO
263 CMO
222 Y1 4-LB
260 CMO
319 PITNSK VC2
342 LITNSOY CMB
238 SHLSCTIVGIIIV TNR-11b
291 TNR-6a
163 TNR-6b

404 LEDIEEALCGPAALP TNR1
431 RSOLETPETLLGSTEER TNR2
401 LLAALRRRIORAD TNR
386 PGSELSTPYQEDGRAM LBR
312 TIILDITSD-SENSWR PMS
246 P COT
530 HKADTVIVGTVKAELPEGROLAGPAELEELEADHTPM CMO
263 QEDGRES CMO
224 FK--OPFMRPVOTT 4-LB
260 RTP CMO
345 PTRF VC2
351 PTHF CMB
251 LIVLLIV TNR-11b
291 SVRER TNR-6a
163 VAGPS TNR-6b

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Figure 4 (continued)

449	P	-----	LPLGVDPACNEPS	-----	PAPSLR	TRF1
449	P	-----	LPLGVDPACNEPS	-----	PAPSLR	TRF2
413	L	VESLCS	-----	-----	TATSPV	TRF
603	M	-----	LAET-ETLGCDDL	-----	NEIOSLV	LTR
139	-----	-----	-----	-----	EPACSP	PS
209	IOED-YRKP	-----	-----	-----	EPACSP	CD7
560	YPEQETEPPLGSCSDVMSVEEKGKEDPLPTAASGK	-----	-----	-----	EPACSP	CD8
270	R	-----	ISVQ-ERQ	-----	GCEL	CD40
236	QEDDC	-----	SC-RFPEEEG	-----	STLAKI	4-1B
283	IQEQADAH	-----	-----	-----	STLAKI	CD40
349	-----	-----	-----	-----	STLAKI	VC22
355	-----	-----	-----	-----	STLAKI	CD48
258	-----	-----	-----	-----	STLAKI	TRF-11be
296	F	-----	-----	-----	STLAKI	TRF-6a
168	L	-----	-----	-----	STLAKI	TRF-6b

Figure 5

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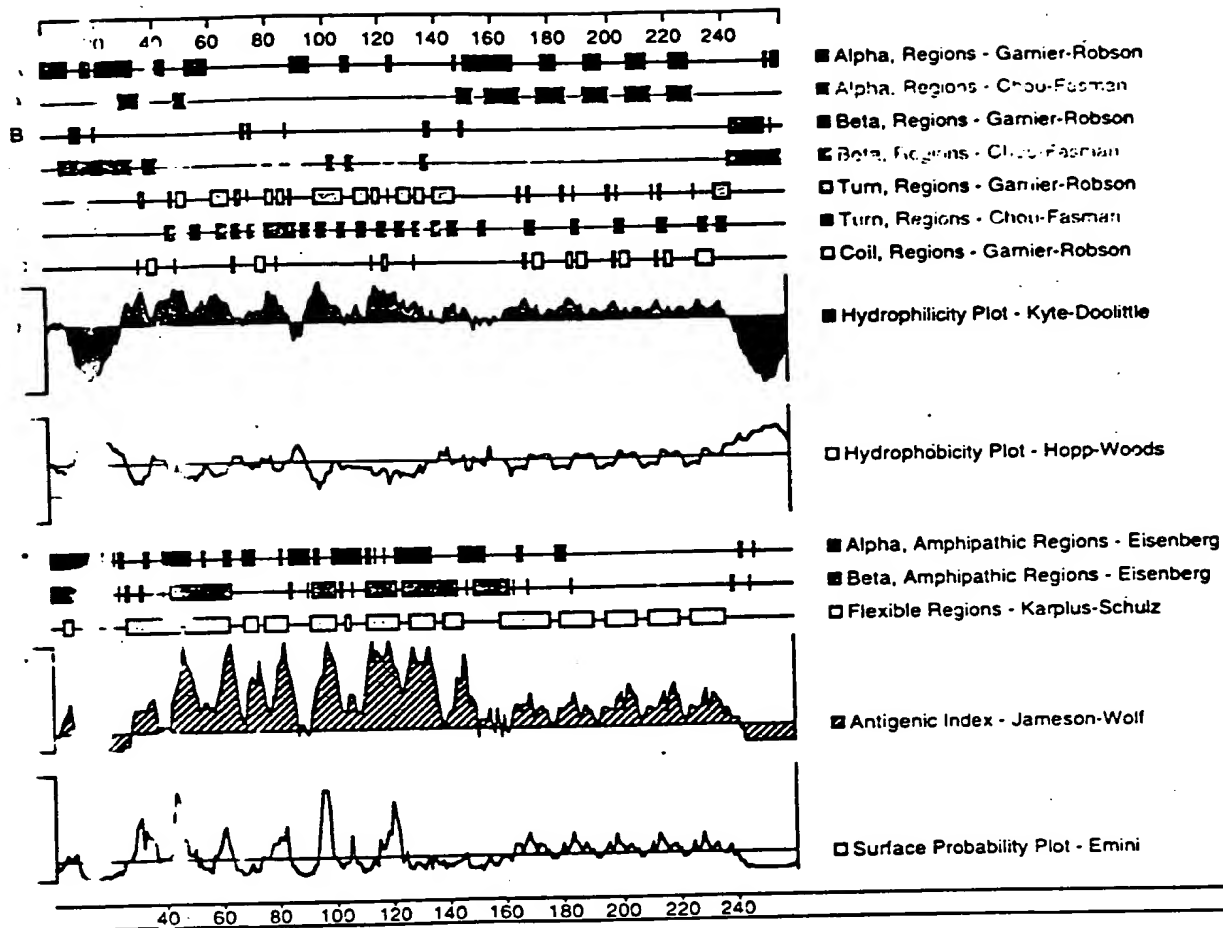


Figure 6

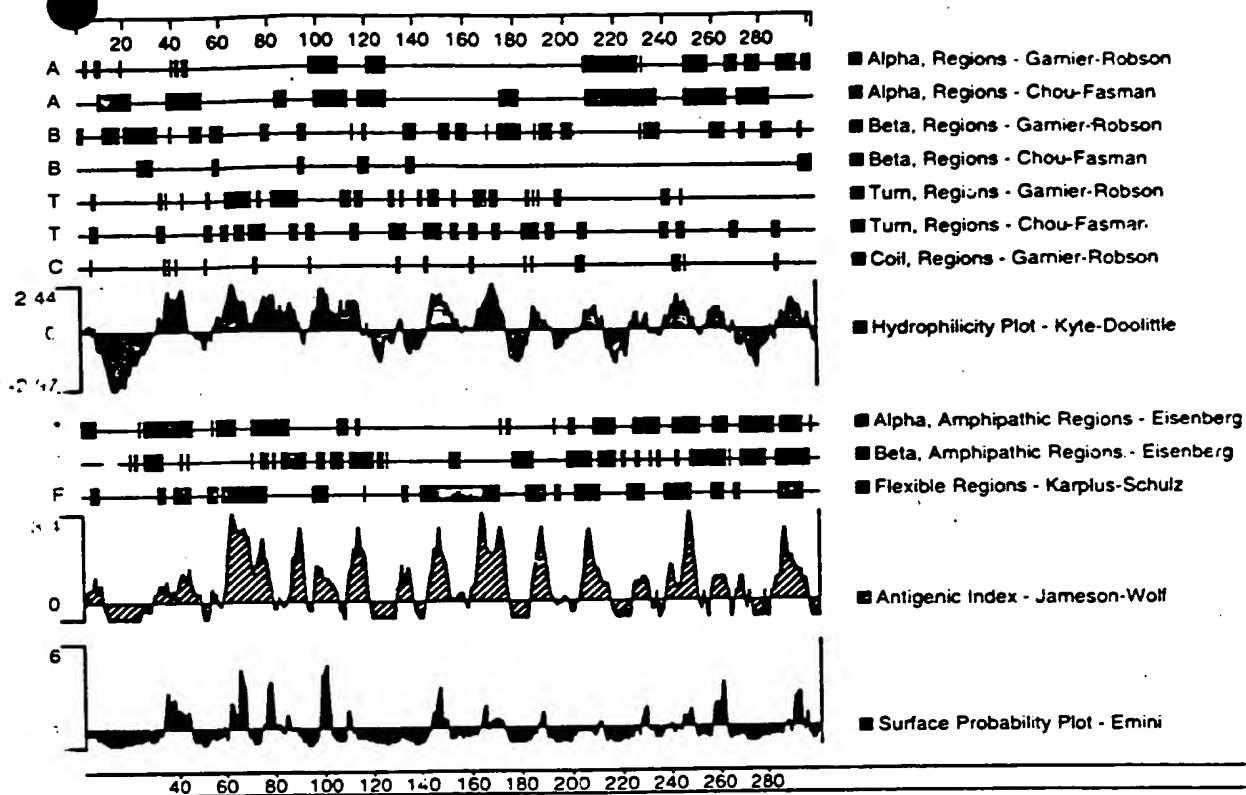




Figure 7

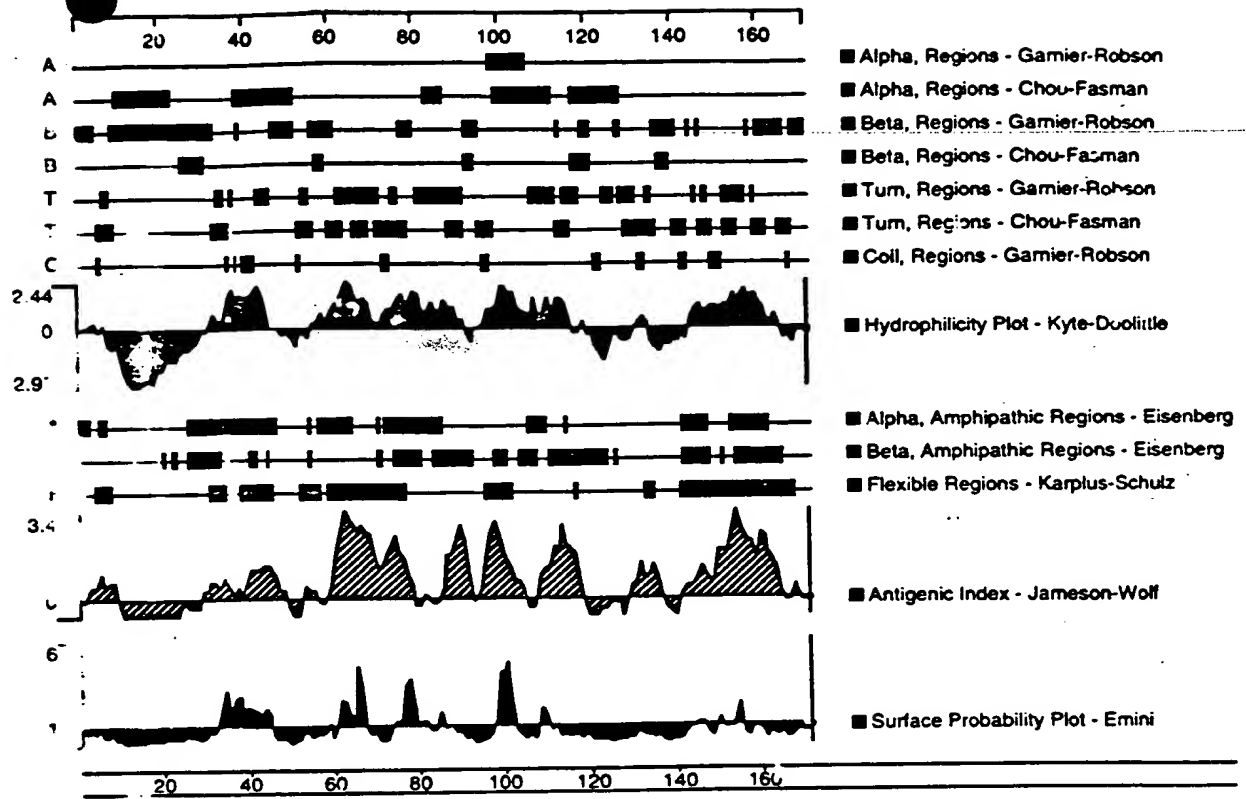


Figure 8

**HELD106R**

GGCAGGAGCA GGGTCCTGTN TCCGCCCTGA GCCGCGCTCT NCCTGCTCCA GCAAGGACCA  
 TGAGGGCGCT GGAGGGGCCA GGCCTGTGCG TGCTGTGCGT GGTGTTGGCG CTGCCTGCCC  
 TGCTGCCGGT GCCGCTGTGA CGCGGAGTGG CAGAAACACN NACNTACCCC TGGCGGGACG  
 NAGAGACAGG GGAGCGGCTG GTGTNTNCCC ANTGCCCCC AGGCACCTTT NTGACGCGG  
 GGTGCCGNG AGACACCCFC ACGACGTGTG GCCGTNTCC ACCGCGCCAC TACACGCATT  
 TGGAACTAC CTGGAGCCTT GNCGTACTN CFACGTCCTG TCGGGGGAGC CTNAGGAGGA  
 GGCACGGGTT TNCCACGNCA ACCACAACCG NGGNTTACCG TNGCCGNACC GGTTCCTTCG  
 TGGCAAGTTG GTTTTNTNTT TGGAGNAAGG ATTCGTGTTN CAATTNATTG ACGNAGTGAT  
 TNNNCNCGGG AAACNTAAA

**HCEOW38R**

CAACTGCA CGGCCCTGGG ACTGGCCCTC AATGTGCCAG GNTCTTCCTC CCATGACACC  
 CTGTGCACCA GCTGCACTGG CTTCCTCCCTC AGCACCAGGG TACCANGAGC TGAGGAGTGT  
 AGCNTGCCG TCATCGACTT TTGGCTTTC CAGGACATCT CCATCAAGAG GCTGCAGCGG  
 GCTCANGC C

**HPRCB54R**

AATTCGGCA NAGCCTCTCC ACGCGCACGA ACTCAGCCAA CGATTTCTGA TAGATTTTGT  
 AGTTTGAC CAGAGATGCA AGGGGTGAAG GAGCGCTTCC TACCGTTAGG AACTCTGGGG  
 ACAGTNGCC CGGCCCGCT GATGGCCGAG GCAGGGTGCG ACCCAGGACC CAGGACGGCG  
 TCGGGAACCA TACCATGGCC CGGATCCCCA AGACCCTAAA GTTCGTGGTC GTCATCGTCG  
 GTCTGTCT GCCAGTCTTA GCTTACTCTG CCACCACTGC CCGSCAGAGG AAGTTNCCCA  
 CAGNANTG GNCCCACAGC AACAGNGSCA CAGTTTCAAG GGGGNAGGAG TTTCCANC  
 TTTTATATA GTTCAGAACN TATTGNGCT NTNAACCTT GCACAAGGGT TTGGNTTAAA  
 CAANGTTTC CAANATGNAC TTTTNGTTC CCTGTTANAT TTTTAATTA GTTNAAANTT  
 AATTTNTNA ACCTTNCCNG GGNAAATT

**H5JA057RA**

GCCAGAGGTG TCTCCAGCCT GGCTCTATCT TCCTCCTTGT NATCGTCCCA TCCCCACATC  
 GTGCAACC CCCAGGACCC TGGTCTCATC AGTCCCTCTC CTGGAGCTGG GGGTCCACAC  
 TCTCCAGC CAAGTCCAAG AGGGCAGGGC CAGTTCTCTC CATCTTCAGG CCCAGCCAGG  
 AGGGGGCAG TCGGCTCCTC AACTGGGTGA CAAGGGTGAG GATGAGAAGT GGTACGGGG  
 TTTTATTCAG CCTTGGTCAG AGCAGAACAC AGAGTTTTTC CGTGTGTTGG TTTTACTCT  
 TTTCCCTT CTTNATNCCC CTTTCN

**HELBP70R**

GCAGAGGCC CCAGCTGCTG AAGAGACAAT AATCACCAGC CCGGGGACTC CTGNTCTNC  
 NATTACCTC TNATGCACCA TCGTAGGGAT CATAGTTCTA ATTGTGCCCT CTAATTGTTT  
 TGTGTTGAAA AGANTTCACT GTGGAAGAAA TTCTTCTCTT ACCTGTAAGT TNCAGGTAGG  
 NGCCTGGCTG AGGGCGGGGG GCGCTGGTAC ACTCTCTGAC CCTGCCTCCC TCTGNTCTGT  
 TCCACAGA CAGAAACGCC TGCCCTGNC CCCAAGTTCC TNGTGTTC CAGCCTGGCT  
 TATCTTNC TCCTGTGAA TCGTTCCAT CCCCACANGC

**HUSCB54R**

GCAGGGTCTC CTNCCCCACC TGCTGAAGAG ACANTGACCA CCAGCCCCGG GACTCCTGCC  
 TCTTCTCAT TACCTCTNAT GNANCATCGT AGGGATCATA GTTCTAATTG TGCCTTCTGA  
 ATTGTGCTTT GTTTGGAAAG ACTTCACTGT GGGGAAGAAAT TCCTTCTT CCGTGAAGTTG  
 CAGGTAGGCC CTGGGTNAGG GCGNGGGCG CTGGACANTN TCTGGNCCTG GCTGCCCCGT